ROCKY FLATS ENVIRONMENTAL TECHNOLOGY SITE

Review of Historical Knowledge Related to Metals and Select Radionuclides Identified as Environmental Media Analytes of Interest

White Paper

Rocky Flats Environmental Technology Site 10808 Highway 93 Golden, CO 80403-8200

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Appendix A

Occurrence Reports

ACRONYMS AND ABBREVIATIONS

AEU Aquatic Exposure Unit

action level AL

AOI analyte of interest

Buffer Zone BZCAA Clean Air Act

CDH Colorado Department of Health

Colorado Department of Public Health and Environment CDPHE

CERCLA Comprehensive Environmental Response, Compensation, and Liability Act

CESC Citizens Environmental Sampling Committee

Colorado Hazardous Waste Act **CHWA** Comprehensive Risk Assessment CRA DOE

U.S. Department of Energy

EPA Environmental Protection Agency

Environmental Restoration ER

EU **Exposure Unit** GW Groundwater

HAER Historic American Engineering Record

hydrochloric acid HC1

HEPA high-efficiency particulate air Historical Release Report HRR

IA Industrial Area

IABZSAP Industrial Area and Buffer Zone Sampling and Analysis Plan

IHSS Individual Hazardous Substance Site IM/IRA Interim Measure/Interim Remedial Action

K-H Kaiser-Hill Company, L.L.C.

kilogram kg

micrograms per liter ug/L mg/day milligrams per day milligrams per kilogram mg/kg

Not Applicable NA ND Not Detected

NESHAP National Emission Standard for Hazardous Air Pollutants

NLR No Longer Representative

Occurrence Reporting and Processing System ORPS

Potential Area of Concern PAC picocuries per gram pCi/g

PCOC potential contaminant of concern **Pre-Demolition Survey Report PDSR**

parts per million ppm

PRG preliminary remediation goal Research and Development R&D

RCRA Resource Conservation and Recovery Act July 15, 2005

RFCA Rocky Flats Cleanup Agreement

RfD reference dose

RFETS Rocky Flats Environmental Technology Site

RFO Rocky Flats Office

RI/FS Remedial Investigation/Feasibility Study

RLCR Reconnaissance-Level Characterization Report

RSOP RFCA Standard Operating Protocol

SAP Sampling and Analysis Plan SEPs Solar Evaporation Ponds UBC Under Building Contamination

UST underground storage tank
VOC volatile organic compound

WEMS Waste and Environmental Management System

WRW wildlife refuge worker

WSRIC Waste Stream and Residue Identification and Characterization Reports

ZPPR Zero Power Plutonium Reactor

1.0 INTRODUCTION

This paper presents a review of information to support a weight-of-evidence determination toward eliminating certain analytes of interest (AOIs) in the Comprehensive Risk Assessment (CRA) potential contaminant of concern (PCOC) professional judgment screening step, on a Sitewide or Exposure Unit (EU)/Aquatic Exposure Unit (AEU) basis. This paper will also be used to supplement evaluations in the Groundwater Interim Measure/Interim Remedial Action (GW IM/IRA) and the nature and extent of contamination sections of the Remedial Investigation /Feasibility Study (RI/FS) Report. The decision to eliminate an AOI from further evaluation will be decided in the GW IM/IRA, and in the specific portions of the RI/FS Report such as the CRA and the nature and extent sections. The information in this paper will be used to supplement the professional judgment decisions within each of these decision documents.

Twenty-four AOIs have been identified in the soil and groundwater nature and extent of contamination evaluations (preliminary reviews), and in sediment and surface water, above preliminary remediation goals (PRGs)¹. These AOIs are composed of 20 metals (other than beryllium, which was used extensively at the Rocky Flats Environmental Technology Site [RFETS]) and four radionuclides (other than americium, plutonium, and uranium isotopes, which were used extensively at RFETS), as listed in Table 1. The reasonably anticipated future land use is a wildlife refuge, therefore Wildlife Refuge Worker (WRW) PRGs were developed using a Hazard Quotient of 0.1 or risk of 1x10⁻⁶ (The more conservative of the two values were used for the PRG). The WRW PRGs were developed specifically for the CRA and are identified in the CRA Methodology (DOE 2004), which has been approved by the Environmental Protection Agency, Region VIII (EPA) and the Colorado Department of Public Health and Environment (CDPHE). WRW PRGs and professional judgment are screening tools used in the CRA and in the RI/FS Report.

Table 2 indicates whether an AOI is addressed in the following sources of information that were reviewed:

- Health Studies on RFETS Phase I: Historical Public Exposures, conducted by ChemRisk for the Colorado Department of Health (CDH) (an independent investigation of off-site health risks associated with operations at RFETS):
 - Project Task 1 Report: Identification of Chemicals and Radionuclides Used at Rocky Flats, March 1991,
 - Project Task 2 Report: Selection of the Chemicals and Radionuclides of Concern, June 1991.
 - Project Tasks 3 & 4 Report: Reconstruction of Historical Rocky Flats Operations & Identification of Release Points, August 1992, and
 - Project Task 5 Report: Estimating Historical Emissions from Rocky Flats 1952-1989;

Building Rocky Flats Cleanup Agreement (RFCA) Reconnaissance Level Characterization Reports (RLCRs) and Pre-Demolition Survey Reports (PDSRs);

¹ The list of surface water AOIs will be updated after a comparison to surface water standards is performed in the surface water nature and extent of contamination evaluation.

Table 1 Preliminary AOIs by Medium

S	oil ^a	Sediment ^a	Surface Water ^a	Groundwater	b	AOIs Evaluated
Surface Soil	Subsurface Soil			UHSU	LHSU	7
Metals	***					
Aluminum		Aluminum				Aluminum
Antimony		Antimony				Antimony
Arsenic	Arsenic	Arsenic	Arsenic	·		Arsenic
						Barium
Cadmium						Cadmium
Chromium	Chromium	Chromium		Chromium (total)		Chromium
Cobalt						Cobalt
						Copper
Iron		Iron				Iron
	Lead					Lead
						Lithium
Manganese		Manganese		Manganese (dissolved)		Manganese
Mercury						Mercury
•						Molybdenum
				Nickel (dissolved)		Nickel
				Selenium (dissolved)		Selenium
						Strontium
			Thallium			Thallium
Vanadium						Vanadium
						Zinc
Radionuclides						
Cesium-137		Cesium-137				Cesium-137
						Strontium-89/90
-		Radium-226		Radium-226 +		Radium-226
Radium-228	Radium-228	Radium-228		Radium-228 (total)		Radium-228

Note: Although barium, lithium, molybdenum, selenium, strontium and zinc (in *italics* in the last column) are not identified as an AOI in a specific medium, they have been identified in the CRA process and as a result are included in this report.

^a AOIs in surface soil, subsurface soil (based on June 30, 2005 draft report), surface water and sediment are those analytes present above a 1x10⁻⁶ WRW PRG.

^b AOIs in groundwater are those analytes present above either a surface water standard or an MCL and form a contiguous plume (based on June 9, 2005 draft report).

Table 2 AOI Usage at RFETS

AOIs	Inventory in RFETS Buildings ^a	Waste Generated in Buildings ^b	Spills/ Releases Within Buildings ^c	Spill/Release Required Action Prior to Demolition ^d	IHSSs/PACs/UBCs Requiring an Accelerated Action ^e	Material of Concern as Selected by ChemRisk Task 2	Material of Concern as Selected by ChemRisk Tasks 3&4
Aluminum	X	X					
Antimony	X	X					
Arsenic	X	X			X		
Barium	X	X					
Cadmium	X	X	X			X	
Chromium (total)	X	X	X		X	X	
Cobalt	X						
Copper	X	X					
Iron	X						
Lead	X	X	X		X	X	
Lithium	X	X	X				
Manganese	X						
Mercury	X	X	X			X	
Molybdenum	X	X	:				
Nickel	X	X	X			X	
Selenium	X	X					
Strontium	X	1338.000					
Thallium	X	X					
Vanadium	X	X					
Zinc	X	X					
Cesium-137	X	X					
Radium-226	X	X					
Radium-228							
Strontium-89/90	X	X					
·	Thorium-232 ^f	X				X	X

^a Based on the ChemRisk Task 1 Report (CDH 1991) and on historical information summarized in the IABZSAP (DOE 2004).
^b Based on the RFETS RCRA Permit, WSRIC, and WEMS (also includes whether underlying hazardous constituents were identified) (See Table 4).

Based on information found in RLCRs and PDSRs (see Tables 3 and 4 for details).

d Sampling of building materials prior to demolition indicated all metal concentrations were below regulatory limits and did not require decontamination or removal.

^eBased on SAPs, SAP Addenda or Closeout Reports for specific IHSS Groups (See Table 5 for details).

f AOI radium-228 is in the thorium-232 decay chain.

- RFETS Resource Conservation and Recovery Act (RCRA)/Colorado Hazardous Waste Act (CHWA) Facility Permit, the Waste Stream and Residue Identification and Characterization (WSRIC) reports, and the Waste and Environmental Management System (WEMS) reports;
- Industrial Area (IA) and Buffer Zone (BZ) SAP (IABZ SAP) Appendix C (to supplement ChemRisk Reports); and
- RFCA Accelerated Action Sampling and Analysis Plans (SAPs), SAP Addenda, and Closeout Reports.

1.1 ChemRisk Tasks 1 through 5 Reports

ChemRisk conducted an independent 2-year investigation of off-site health risks associated with operations of the Rocky Flats Plant for the CDH (Tasks 1 through 12). This investigation generated an inventory of chemicals and radionuclides that have been used or produced at RFETS. This information was screened in various Task Reports to identify a "short list" of chemicals that, because of the amounts, processes, and duration of use, should be evaluated for off-site release potential.

The buildings² identified in the Task 3 & 4 ChemRisk report formed the basis for evaluating the usage of the AOIs in this review.

ChemRisk Task 1 involved the identification of chemicals and radionuclides that have been used on-site. For these chemicals, a three stage screening process was developed to narrow down the list of potential materials of concern. Initially, over 8,000 chemicals were identified in the Task 1 Report. Screening stages were developed in the Task 2 Report to help evaluate the list of chemicals, based on such factors as the relative toxicity of the materials, quantities used, how the materials might have been released into the environment, and the likelihood for transport of the materials off-site. In the first stage, 629 compounds were identified for further, more refined screening as potential materials of concern (as defined by the ChemRisk process based on materials in inventory, which may pose an off-site health risk) based on their known toxicologic properties, RFETS release histories, or reported inventory quantities. (Material of concern is defined by the ChemRisk process as the inventory of materials used at RFETS, which could pose an off-site health risk.) A second stage of screening was performed to roughly estimate if the quantity of a chemical on-site was sufficient to pose an off-site health hazard. Forty-six potential chemicals of concern emerged from Stage 2 screening. In the final stage of screening, these chemicals were individually evaluated to determine the likelihood of their release and potential quantity of release based on actual storage and usage practices, likely routes of release, and known behavior in the environment. Based on this final screen a total of 25 materials of concern were identified in the Task 2 Report and further evaluated in the Task 3 & 4 Report. Of these 25, only five metals (Table 2) were identified and eventually dropped in the Task 3&4 Report.

² The buildings discussed in the ChemRisk reports are presented later in this report in Tables 3 and 4.

1.2 RLCRs and PDSRs

To supplement the historic usage information, building specific information from RLCRs and PDSRs were used to identify any contaminants that may have been present in buildings (discussed in the ChemRisk Reports) prior to demolition (including spills or releases) or, in the case of Type 2 or 3 buildings, were the basis of a hazard profile analysis for a building (Table 3). A building has never been classified as either a Type 2 or 3 building based on the presence of any of the AOI metals or radionuclides. The Type 2 and 3 classifications were primarily based on the presence of beryllium and/or americium, plutonium, or Uranium radionuclides. In addition, all RCRA units within buildings were either certified clean-closed or were closed by removal prior to demolition. (Additional information is presented below regarding the history of RCRA units.)

1.3 RCRA/CHWA Facility Permit, WSRIC Reports and WEMS Reports

Hazardous waste (including mixed waste) management activities were conducted in many buildings at RFETS. Table 4 lists the hazardous wastes managed in buildings (discussed in the ChemRisk Reports). A review was conducted of the RFETS RCRA/CHWA Operating Permit, as well as the waste³ generated from the various buildings as identified in the WSRIC Reports and the WEMS Reports (part of the RCRA operating record used to track and control inventory and movement of hazardous, nonhazardous, and mixed waste containers). Based on process knowledge, RCRA waste codes were conservatively applied to wastes generated within the buildings. For example, if the possibility existed for one building to generate a RCRA waste, all process buildings were identified to also carry this waste code in case waste was transferred via process waste lines or moved into a separate building for storage and/or treatment or if analysis was required on the waste. Also, historically, RCRA waste codes were conservatively applied to materials without specific attention given to concentration (when mixtures were involved) or to the process generating the waste.

Specific units were permitted under the RFETS RCRA/CHWA Operating Permit or were operated under RCRA/CHWA Interim Status requirements, pending their closure. Closure of permitted and interim status units is governed under the Colorado Department of Public Health and Environment (CDPHE) approved closure plans. The focus of closure plans is to ensure that any hazardous wastes in the unit have been removed and that any unit components with residual wastes are properly decontaminated or removed, resulting in unit clean closure. Documentation that wastes have been removed and all necessary decontamination and component removal have been conducted was submitted to CDPHE for approval of the unit closure. While spills related to some units may have occurred during operations, the permit and interim status requirements governed the appropriate cleanup response action, including decontamination if necessary, taken at the time. These actions prevented any significant impacts by prompt and effective removal of spilled hazardous wastes.

The RLCRs include information on hazardous waste units, characterization of residual hazardous wastes, and unit closure. Inspection and characterization of these units indicated residual hazardous waste contamination was basically confined to unit components, such as tanks, piping, floors, and floor coverings and sumps designed to contain these wastes during waste management activities. The units were properly closed prior to building decommissioning in accordance with

³ This review was conducted based on RCRA waste codes and included all waste that may have carried a specific waste code. This review also looked at any waste associated with non-RCRA metals such as aluminum.

Table 3 History of AOIs Used in RFETS Buildings

AOIs	371 a,b,d	374 ^{a,b,d}	444 ^{a,b}	445 ^{b,c}	450 ^{b,c}	455 ^{b,c}	460 ^{b,d}	559 ^{a,b} (Lab)	561 ^{b,c}	707 ^{a,b}	712/ 713 ^{b,c}	774 ^{a,b}	771 ^{a,b}	776/ 777 ^{b,c}	779 ^{b,c} (Lab)	881 ^{b,c} (Lab)	883 ^{b,c}	865 ^{b,c}
Metals														r.		r	,	
Aluminum			X							X			X		X		X	X
Antimony ^e																		
Arsenic ^e								•••						, ,				
Barium																		
Cadmium			X					X	X	X		X	X	X		X	X	
Chromium	X	X	X				X	X	X		X	X		X	X			
(total)																		
Cobalte																		L
Copper			X												X	X	X	X
Iron	X		X	X	X	X			ļ									L
Lead	X	X						X		X		X	X	X				
Lithium								X		X				X		X		
Manganese																		
Mercury								X	X					X		X		
Molybdenum			X										X	X				X
Nickel	X	X	X									X	X	X	X	X		
Selenium°																		
Strontium ^e																		
Thallium ^e																		<u> </u>
Vanadium			X							X					X	X	X	X
Zinc									,									
Radionuclides		,																
Cs-137 ^e																		
Ra-226 ^e																		
Ra-228																		
Sr-89/90 ^e															X		· ·	
Thorium-232													X			X		

Note those AOIs identified in *Italic type* above indicate that these were included in the ChemRisk inventories for the Site, however there were no specific building identified. Those AOIs identified in **bold type** indicate RCRA metals.

Footnotes are based on PDSRs, RLCRs and Historical Site Assessment Reports (attached to RLCR).

Lab = Laboratory

- a Spills occurred, areas sampled and results indicate all concentrations below RCRA regulatory limits for RCRA metals.
- b Buildings contained RCRA units which were closed by demonstrating clean closure or closure by removal.
- c No known spills of RCRA/CERCLA contaminants.
- d PDSRs for all phases of work are currently not available.
- e These AOIs were identified in very small quantities associated with laboratory operations and used as laboratory standards or in analytical testing.

Table 4 Waste containing AOI generated in RFETS Buildings

AOIs	371	374	444	445	450	455	460	559	561	707	712/713	771	774	776/777	779	881	883	865
Metals																		
Aluminum	Xª							X		Xª					X			
Antimony								X ^b										
Arsenic	X	X					X	X		X		X	X	X	X	X	X	X
Barium	X	X	Xb					X		X		X	X	X	X	X	X	X
Cadmium	X	X	X				X	X		X		X	X	X	X	X	X	X
Chromium	X	X	X				X	X	•	X		X	X	X	X	$\bar{\mathbf{X}}$	X	X
(total)																		
Cobalt																		
Copper								X								X		
Iron																		
Lead	X	X	X				X	X		X		X	X	X	X	X	X	X
Lithium	X ^c		X°				X ^c	X				X ^c				X	X	
Manganese																		
Mercury	X	X	X				X	X		X		X	X	X	X	X	X	X
Molybdenum								X							<u> </u>			
Nickel	X_p	Xb	Xb					X ^b		Xb		X ^b	Xb	Xb	Xb	Xb	X ^b	X ^b
Selenium	X	X	X					X		X		X	X	X	X	X	X	X
Strontium																		
Thallium			<u> </u>					X ^b		X ^b		X ^b			Xb	Xb		
Vanadium								Xb								Xb		
Zinc				<u> </u>							<u>.</u>					Xb		
Radionuclides																		
Cesium-137			<u> </u>						<u></u> .	Xd				X ^d	<u> </u>			
Radium-226						L				X ^d				X^{d}				
Radium-228																		
Strontium- 89/90										Xd								
Thorium-232	-	idontifia														X^d		

Note those AOIs identified in **bold type** above indicate RCRA metals. Footnotes are based on printouts from WSRIC.

 ^a Aluminum oxide and magnesium oxide crucibles.
 ^b Analyte was identified as an underlying hazardous constituent and not as a RCRA toxicity metal waste.
 ^c Waste consisted of Lithium batteries.
 ^d Sealed sources were removed as waste from this building.

the permit or interim status closure plans, or as part of the decommissioning process under the RFCA decision documents and closures approved by CDPHE. Thus, RCRA/CHWA closure activities did not indicate any significant releases of hazardous wastes from these units. The WEMS database was implemented in 1990 and the WSRIC database in January 2002⁴. Any information regarding wastes and any spills resulting in wastes prior to this time would have been included in the evaluation performed by ChemRisk and by the Environmental Restoration (ER) program.

1.4 IABZ SAP, Appendix C

Appendix C of the IABZ SAP contains historical building process information summarized from the 1998 Historic American Engineering Record (HAER) (for the Rocky Flats Historic District). This information was used to supplement chemical usage information from the ChemRisk reports.

1.5 SAPs, SAP Addenda, and Closeout Reports

A summary is provided in Table 5 based on SAPs, SAP Addenda, and Closeout Reports for Individual Hazardous Substance Sites (IHSSs), Potential Area of Concern (PACs) or Under Building Contamination (UBCs) sites to indicate whether the presence of metals in the environment were identified above RFCA action levels (ALs), and were removed under an IHSS accelerated action.

Table 5 Metals requiring an accelerated action at IHSSs/PACs/UBC Sites

IHSS Group	IHSS/PAC/UBC Site	Building/ Structure	Metal Requiring Action
700-6	PAC 700-137 (Cooling Tower Blowdown)	Buildings 712/713	Arsenic
100-4	UBC 123	Building 123	Lead
400-8	PAC 400-122	UST associated with Building 441	Lead
500-2	PAC 500-158	Building 551	Chromium
700-2	No specific IHSS	Outside of Building 707	Arsenic
NE-1	PAC NW-1505	North Firing Range	Lead
900-11	PAC SE-1602	East Firing Range	Lead and Arsenic

In addition to the buildings identified in the ChemRisk reports and presented in Tables 3 and 4, potential UBC sites (that were not included in the ChemRisk buildings) have also been added and presented in Tables 6 and 7. These buildings were added due to the potential contamination

⁴ Although the WSRIC database was not implemented until 2002, building books were started in 1991 and this information was included in the database.

Table 6 History of AOIs Used in UBC Sites

AOIs	122 ^{c,e}	123 a,b	125 a	331 ^c	439°	440 ^d	441ª	442	447 ^{a,b}	528 ^{a,b}	701 ^d	731 a,b,d	770	778	886 Lab	887 ^{a,b}	889	991 ^{b,c}
Metals		Lab	Lab			L		<u>i</u>	<u> </u>				<u> </u>	1	Lau	1		
Aluminum							Ţ		X				Ī	İ				
Antimony																		
Arsenic																		
Barium																		
Cadmium		X								X		X						
Chromium		X								X		X						
(total)																		
Cobalte																		
Copper																		
Iron																		
Lead		X								X	X	X						
Lithium																		
Manganese																	·	
Mercury															X			
Molybdenum																		
Nickel																		X
Selenium ^e																		
Strontiume																		
Thallium ^e																		
Vanadium									X									
Zinc																		
Radionuclides															"			
Cs-137 ^e															X			
Ra-226 ^e																		
Ra-228																		
Sr-89/90 ^e																		
Thorium-232		OIs identi																

Note those AOIs identified in **bold type** indicate RCRA metals.

Footnotes are based on PDSRs, RLCRs, Closeout Reports and/or Historical Site Assessment. Reports (attached to RLCR).

Lab = Laboratory

- a Spills occurred, areas sampled and results indicate all concentrations below RCRA regulatory limits for RCRA metals.
- b Buildings contained RCRA units which were closed by demonstrating clean closure or closure by removal.
- c No known spills of RCRA/CERCLA contaminants.
- d PDSRs and/or Closeout Reports for all phases of work are currently not available.
- e These AOIs were identified in very small quantities associated with laboratory operations and used as laboratory standards or in analytical testing.

Table 7 Waste containing AOI generated in UBC Sites

AOIs	122	123	125	331	439	440	441	442	447	528	701	731	770	778	886	887	889	991
Metals																		
Aluminum														Xª		Xa		
Antimony																		
Arsenic														X		X	X	
Barium					X			Xb						X	X	X		
Cadmium	X	X	X	X	X	X	Xb	X ^b			X			X	X	X		X
Chromium	Xb	X		X	X	X	X^{b}		X		X			X	X	X		X
(total)																		
Cobalt																		
Copper																		
Iron																		
Lead	X	X	X	X	X	X		X	X		X		X	X	X	X	X	X
Lithium			X ^c											X ^c				
Manganese												ļ. <u>.</u>						
Mercury		X	X	X	X	X		X			X		X	X	X	X	X	X
Molybdenum																		
Nickel	Xb		Xb	Xb	X^{b}			Xb			X_p			X^{b}	Xb			Xb
Selenium					X		X ^b			,						X		
Strontium																		
Thallium																		
Vanadium																		
Zinc											,	<u> </u>						
Radionuclides																		
Cesium-137																		X^{d}
Radium-226																		
Radium-228																		
Strontium-																		X ^d
89/90																		
Thorium-232								L										

Note those AOIs identified in **bold type** above indicate RCRA metals.

Footnotes are based on printouts from WSRIC.

^a Aluminum silicate (clay) identified to be associated with "combustibles contaminated with adhesives".

^b Analyte was identified as an underlying hazardous constituent and not as a RCRA toxicity metal waste.

^c Waste consisted of Lithium batteries.

^d Sealed sources were removed as waste from this building.

of soil identified or suspected under specific buildings from broken process waste lines or other sources. The information used to complete these tables is the same information as described in sections 1.2 through 1.4 and from the site Historical Release Report (HRR) (DOE 1992).

Additional sources of information that were reviewed but did not identify metal concerns are discussed below.

1.6 Compliance Review under the CAA

National Emission Standards for Hazardous Air Pollutants (NESHAPs), developed pursuant to the Clean Air Act (CAA), have not been applicable to any processes on Site involving the usage of metals. In addition, emission calculations for specific processes (for example, Building 443 boiler emissions from fuel consumption) have consistently⁵ indicated metal concentrations are below any reporting threshold.

1.7 ORPS

A review of the Occurrence Reporting and Processing System (ORPS) occurrence reports was conducted (from 1991 to current) for emergencies at RFETS that required implementation of either the Site emergency plan or the Site RCRA contingency plan involving buildings and spill/releases into the environment. For occurrence reporting prior to 1991, no electronic system was available, and lists of occurrences, event reporting, health and safety issues, and serious incidence reports were reviewed instead for the time period between 1952 and 1990. Very few documented incidents (within a span of 50 years) occurred within a building that would have resulted in a release to the environment. Of those incidents that could have impacted the environment, all were historically identified as an IHSS, PAC, or UBC Site and evaluated to determine if an accelerated action was needed. All accelerated actions have resulted in a No Further Accelerated Action (NFAA) determination.

Between 1991 and the present, only three occurrences were reported that potentially involved a release of an AOI outside of a building. These occurrences resulted in either no impact to the environment or no significant impact to the environment.

In 1993 there was a sprinkler head malfunction in Room 3189 (a radiological material area) of Building 374, releasing firewater that eventually ran onto Dock 18T and onto the ground. Sample results determined the water was clean and approved for release to the stormdrain system. There was no impact to the environment (RFO 1993 in Appendix A, Occurrence Reports).

In 1998 approximately 1 gallon of phosphoric acid contaminated with depleted uranium was released from a dock drain line into a bermed area from Tank D-843 at Building 371. The occurrence report indicated there was no impact to the environment (RFO 1998 in Appendix A). Characterization data around the 371 dock did not indicate any uranium concentrations greater than background mean plus two standard deviations, based on the SAP Addendum for IHSS Group 300-3 (DOE 2003a).

⁵ A thorough evaluation of emission inventories began at RFETS in late 1989.

In 2002 there was a spill of low-level mixed waste from a RCRA-regulated tank located south of Buildings 371/374. Approximately 1 to 5 gallons of waste were released outside the secondary containment over an area of approximately 600 square feet. Analytical results for all RCRA metals were below regulatory limits, with cadmium results being the highest at 104 micrograms per liter (µg/L) (parts per billion) (RFO 2002 in Appendix A). This spill was remediated as part of accelerated actions at RFETS.

2.0 SUMMARY OF INFORMATION AVAILABLE FOR AOIS

Based on the various categories of information reviewed (including the ChemRisk reports), a summary is provided below for each of the 23 AOIs.

Figures 1 through 22 provide soil sampling and analysis results for data (June 28, 1991 through April 27, 2005) greater than background mean plus two standard deviations (as defined in Attachment 2, Volume 2 of 15, Appendix A of the RI/FS Report (DOE 2005a) beneath the slabs⁶ at UBC sites and surface soil data around the buildings. This data conservatively represent the contaminants that remain, since more accelerated actions have been completed (after April 27, 2005) at the site resulting in more data becoming no longer representative (NLR).

Tables 8 through 27 provide summary statistics for the UBC soil analytical results, including a comparison to surface soil background mean plus two standard deviations (for Buildings 371, 374, 444, 559, 561, 707, 771, 774, 776, 777, 779, 865, 881, and 883). The remaining buildings for which UBC investigations were conducted have also been included in both figures and tables. These include Buildings 122, 123, 125, 331, 439, 440, 441, 442, 447, 528, 701, 731, 770, 778, 886, 887, 889, and 991. Although Buildings 125, 460, 561, 731 and 887 were included, there is no data reported in Tables 8 through 27 for the following reasons:

- Building 125 received a no further accelerated action determination;
- Building 460 was not expected to have contamination beneath the slab;
- Building 561 was not identified as a UBC building;
- Samples were collected around Building 731 at depths below the slab, but not beneath the slab since groundwater would have been encountered under artesian conditions; and
- Building 887 samples were moved to the outside of the building footprint because of concerns with groundwater beneath the building.

The figures and tables indicate a significant portion of AOI mean concentrations are below background mean plus two standard deviations (antimony, arsenic, barium, cadmium, lead, manganese, mercury, molybdenum, selenium and zinc). For aluminum, chromium, cobalt, iron, lithium, nickel, strontium, and vanadium, the mean concentrations are within or very closely approximate background. In reviewing the figures, the horizontal spatial distribution of AOI concentrations do not indicate large areas of soil with concentrations that exceed background, and point to a very limited surficial soil or UBC deposition. For copper, and thallium a majority

⁷ Each figure identifies concentrations of metals greater than the WRW PRG (1x10⁻⁶ risk level) and 10 times WRW PRG (1x10⁻⁵ risk level).

⁶ Data includes samples at depth intervals of 0 to 0.5 feet beneath the building slabs. Also included were samples collected at a depth interval of 0 to 2.5 feet or 3.0 feet, if this was the uppermost sample interval beneath the slab.

of the mean concentrations beneath UBCs are above background (but below RFCA action levels for accelerated actions).

Based on extensive experience in soil removal at IHSSs, such limited areas of elevated surficial and UBC concentrations indicate subsurface migration of AOIs at higher concentrations would not be expected. Rather, this experience indicates generally that concentrations significantly decrease with depth (metals if soluble are influenced by the chemistry of the soil [for example, pH, the presence of other metals and oxygen] and do not move significantly in the subsurface). It is unlikely that these limited elevated concentration locations result in subsurface soil contamination or present a source of groundwater contamination.

2.1 Metals

A total of 20 metal AOIs have been identified in media based on the nature and extent evaluations. These metals are discussed in the following sections.

2.1.1 Aluminum

Aluminum was used primarily in various metallurgical operations within Buildings 444, 779, 865, and 883 (CDH 1992; DOE 2004). Rejected aluminum parts were disassembled and recycled or prepared for disposal in Building 707. Aluminum was also used in pit construction (Building 707). In Building 447 materials handled included stainless steel, beryllium, aluminum, depleted uranium, and vanadium compounds. Aluminum nitrate was used in an aqueous dissolution process within Building 771 for plutonium recovery.

All of the buildings identified above (except Building 778 [Plant Laundry Facility]) involved radiological operations and included extensive high efficiency particulate air (HEPA) filtration systems. Any particulates or fines from machining aluminum metals would have been collected on these filters prior to release from the buildings. (Building 887 was used to manage process and sanitary waste, and Building 778 laundered clothing. These buildings did not machine aluminum metal.)

Aluminum was identified in the ChemRisk reports as a chemical (for example, aluminum nitrate) and not as a metal (CDH 1991a). Aluminum nitrate was not carried forward as a material of concern for the ChemRisk reports based on ingestion of this material in a drinking water exposure scenario for off-site receptors (CDH 1991b).

There is no record of spills involving aluminum nitrate within these buildings, based on a review of RLCRs and PDSRs for these buildings. In the remaining UBC buildings, only aluminum silicate (clay) was identified as being used in Buildings 778 and 887.

Six buildings identified generating aluminum waste. Two buildings (371 and 707) identified the waste to be aluminum oxide and magnesium oxide crucibles. Two buildings (778 and 887) identified the waste to be aluminum silicate (clay) combustibles associated with adhesives. Building 779 waste was identified as a desiccant aluminum oxide and Building 559 waste was identified as processing agents that may contain lithium aluminum hydride.

Aluminum was not identified above a RFCA AL requiring an accelerated action based on SAPs, SAP Addenda, or Closeout Reports for IHSSs and UBC sites.

In reviewing the aluminum soil data beneath the slabs for UBC sites (Figure 1) the summary statistics presented in Table 8 were generated.

Table 8 UBC Soil Summary Statistics for Aluminum (mg/kg)

	1				1	(8/8/	· · · · · · ·	·
UBC	Analyte	Total Number of Samples	Detection Frequency > Background ⁸	Mean	Maximum Concentration	Standard Deviation	Background Mean Plus 2SD	Unit
122	Aluminum	6	33.33%	10,550	19000	7,879	16,715	mg/kg
123	Aluminum	0	NA	_ NA	NA	NA	16,715	mg/kg
331	Aluminum	16	56.25%	15,900	22000	3,871	16,715	mg/kg
371	Aluminum	1	0.00%	2,200	2200	NA	16,715	mg/kg
374	Aluminum	3	0.00%	4,123	6870	2,600	16,715	mg/kg
439	Aluminum	6	0.00%	6,585	11000	2,967	16,715	mg/kg
440	Aluminum	10	20.00%	11,610	20000	5,514	16,715	mg/kg
441	Aluminum	5	100.00%	22,000	27000	3,873	16,715	mg/kg
442	Aluminum	11	45.45%	17,403	29000	7,224	16,715	mg/kg
444	Aluminum	39	43.59%	15,846	31000	6,037	16,715	mg/kg
445	Aluminum	1	100.00%	37,000	37000	NA	16,715	mg/kg
447	Aluminum	17	88.24%	22,818	38000	7,467	16,715	mg/kg
450	Aluminum	1	100.00%	17,000	17000	NA	16,715	mg/kg
455	Aluminum	1	0.00%	6,600	6600	NA	16,715	mg/kg
528	Aluminum	1	100.00%	20,000	20000	NA	16,715	mg/kg
559	Aluminum	12	16.67%	12,333	22000	5,533	16,715	mg/kg
701	Aluminum	4	50.00%	15,200	24000	6,814	16,715	mg/kg
707	Aluminum	20	0.00%	5,390	13000	2,406	16,715	mg/kg
712/713	Aluminum	12	50.00%	18,558	44000	12,452	16,715	mg/kg
770	Aluminum	2	50.00%	16,500	18000	2,121	16,715	mg/kg
771	Aluminum	12	8.33%	13,131	21000	3,704	16,715	mg/kg
774	Aluminum	4	100.00%	18,750	21000	1,500	16,715	mg/kg
776	Aluminum	20	35.00%	13,875	33000	8,745	16,715	mg/kg
777	Aluminum	25	8.00%	10,504	35000	7,334	16,715	mg/kg
778	Aluminum	15	13.33%	11,040	21000	5,778	16,715	mg/kg
779	Aluminum	3	100.00%	29,667	32000	2,082	16,715	mg/kg
865	Aluminum	25	28.00%	13,308	36000	8,184	16,715	mg/kg
881	Aluminum	25	28.00%	15,800	23000	3,765	16,715	mg/kg
883	Aluminum	12	16.67%	12,908	22000	4,397	16,715	mg/kg
886	Aluminum	23	21.74%	12,684	59000	11,910	16,715	mg/kg
889	Aluminum	3	0.00%	14,433	16300	1,665	16,715	mg/kg
991	Aluminum	10	60.00%	16,040	37000	10,597	16,715	mg/kg
- N.T.A	- Not Appli	1.1.			·	.		

NA = Not Applicable

⁸ This column indicates those samples that were detected and greater than background. As a result, if this field shows 0% and a mean and maximum concentration are reported, the values represent those analytes detected but less than background.

2.1.2 Antimony

Antimony was not identified or discussed in building process information (CDH 1992; DOE 2004). Antimony has not been found associated with UBC sites (DOE 2004).

Antimony was initially identified in the ChemRisk Task 1 Report as a chemical in inventory at RFETS (although no specific building was identified). Examples include antimony in 20% hydrochloric acid (HCl) solution, antimony iodide, antimony oxide, antimony pentachloride, antimony powder, antimony trioxide, and antimony trichloride. These chemicals appeared to have been used as laboratory standards or analytical testing materials because they were used in very small quantities. This is confirmed based on a review of waste generated within process buildings, where antimony was identified to be present within only one RFETS building (559), which was a laboratory building (Table 4). Antimony was not carried forward as a material of concern for the ChemRisk process indicating an insufficient quantity existed at RFETS to pose a potential off-site health hazard (CDH 1991a, 1991b).

Building 559 is the only building identifying waste containing antimony and as an underlying hazardous constituent.

At the bermed area east of the North Target Area at the East Firing Range, one sample out of thirty (CW37-012) identified antimony at a concentration (433 mg/kg) greater than the RFCA AL (409 mg/kg) (DOE 2005b). This surface soil was removed (DOE 2005). Antimony was not identified at other areas of the site above a RFCA AL requiring an accelerated action based on SAPs, SAP Addenda, or Closeout Reports for IHSSs and UBCs.

In reviewing the antimony soil data beneath the slabs for UBC sites (Figure 2), the summary statistics presented in Table 9 were generated.

		abic 7 CDC	Son Summar	y Statistics it	n Anumony (ing/kg)		
UBC	Analyte	Total Number of Samples	Detection Frequency > Background ⁹	Mean	Maximum Concentration	Standard Deviation	Background Mean Plus 2SD	Unit
122	Antimony	6	0.00%	0.167	0.3	0.065	0.436	mg/kg
123	Antimony	0	NA	NA	NA	NA	0.436	mg/kg
331	Antimony	16	31.25%	0.382	0.77	0.219	0.436	mg/kg
371	Antimony	1	0.00%	ND	ND	ND	0.436	mg/kg
374	Antimony	3	0.00%	ND	ND	ND	0.436	mg/kg
439	Antimony	6	0.00%	ND	ND	ND	0.436	mg/kg
440	Antimony	10	30.00%	0.361	1.4	0.420	0.436	mg/kg
441	Antimony	5	0.00%	ND	ND	ND	0.436	mg/kg
442	Antimony	11	9.09%	0.257	0.58	0.107	0.436	mg/kg
444	Antimony	39	7.69%	0.192	0.57	0.112	0.436	mg/kg
445	Antimony	1	0.00%	ND	ND	ND	0.436	mg/kg
447	Antimony	17	17.65%	0.280	0.46	0.132	0.436	mg/kg
450	Antimony	1	0.00%	ND	ND	ND	0.436	mg/kg
455	Antimony	1	0.00%	ND	ND	ND	0.436	mg/kg

Table 9 UBC Soil Summary Statistics for Antimony (mg/kg)

⁹ This column indicates those samples that were detected and greater than background. As a result, if this field shows 0% and a mean and maximum concentration are reported, the values represent those analytes detected but less than background.

				<u> </u>				
UBC	Analyte	Total Number of Samples	Detection Frequency > Background9	Mean	Maximum Concentration	Standard Deviation	Background Mean Plus 2SD	Unit
528	Antimony	1	0.00%	ND	ND	ND	0.436	mg/kg
559	Antimony	12	25.00%	0.279	0.58	0.184	0.436	mg/kg
701	Antimony	4	0.00%	ND	ND	ND	0.436	mg/kg
707	Antimony	20	0.00%	0.163	0.43	0.072	0.436	mg/kg
712/713	Antimony	12	33.33%	0.934	4	1.26	0.436	mg/kg
770	Antimony	2	0.00%	ND	ND	ND	0.436	mg/kg
771	Antimony	12	0.00%	ND	ND	ND	0.436	mg/kg
774	Antimony	4	0.00%	ND	ND	ND	0.436	mg/kg
776	Antimony	20	5.00%	0.167	0.44	0.076	0.436	mg/kg
777	Antimony	25	0.00%	0.158	0.3	0.051	0.436	mg/kg
778	Antimony	15	13.33%	0.281	1.1	0.308	0.436	mg/kg
779	Antimony	3	0.00%	0.213	0.33	0.101	0.436	mg/kg
865	Antimony	25	0.00%	0.153	0.38	0.049	0.436	mg/kg
881	Antimony	25	24.00%	0.307	0.6	0.139	0.436	mg/kg
883	Antimony	12	25.00%	0.354	1.2	0.300	0.436	mg/kg
886	Antimony	22	9.09%	0.202	0.59	0.128	0.436	mg/kg
889	Antimony	3	0.00%	ND	ND	ND	0.436	mg/kg
991	Antimony	10	0.00%	0.172	0.36	0.067	0.436	mg/kg

NA = Not Applicable

ND = Not Detected

2.1.3 Arsenic

Arsenic was not identified or discussed in building process information (CDH 1992; DOE 2004). Arsenic has not been found associated with UBC sites (DOE 2004).

Arsenic was initially identified in the ChemRisk Task 1 Report as a chemical in inventory at RFETS (although no specific building was identified) as well as a likely organic-arsenical compound found in pesticides used at RFETS. Examples include arsenic acid, arsenic iodide, arsenic metals, arsenic pentoxide, arsenic solution 3103, arsenic trioxide, arsenious oxide, and arsenious acid (CDH 1991a). These chemicals were identified to be present at RFETS in very small quantities (less than 1 kilogram [kg]), and were identified as laboratory standards used in Buildings 444, 559, 779, and 881. The Task 2 report concluded that based on the limited use of these chemicals and their annual usage rates, which were greater than inventory quantities, their release to the environment was estimated to be minimal or there would be no release (CDH 1991b).

Arsenic waste has been generated from both laboratory and process buildings (Table 4), and at Building 778 (Plant Laundry Facility), Building 887 (managed process and sanitary waste), and Building 889 (Decontamination and Waste Reduction Facility) (Table 7).

There is no record of spills involving arsenic within these buildings, based on a review of RLCRs, PDSRs and Closeout Reports for these buildings.

Arsenic was identified as present in soil above the RFCA AL requiring an accelerated action based on SAPs, SAP Addenda, or Closeout Reports for IHSSs and UBC sites. Specifically, at

the Building 712/713 cooling towers (IHSS 700-137), in which arsenic may have been a component of the rust inhibitors used in the cooling towers, and at the East Firing Range (IHSS SE-1602) as a component in lead shot. In addition to these two areas, arsenic was also identified to be present at each of the downspouts to Building 707 (IHSS Group 700-2) (at concentrations above the RFCA AL), which may have been associated with rat poison used on the roof or the presence of treated lumber also located on the roof.

In reviewing the arsenic soil data beneath the slabs for UBC sites (Figure 3), the summary statistics presented in Table 10 were generated.

Table 10 UBC Soil Summary Statistics for Arsenic (mg/kg)

	1 4	DIC TO ODC	Son Summary	Statistica	TOI AISCINC	(mg/Kg)		
UBC	Analyte	Total Number of Samples	Detection Frequency > Background ¹⁰	Mean	Maximum Concentration	Standard Deviation	Background Mean Plus 2SD	Unit
122	Arsenic	6	0.00%	4.42	9.1	3.50	10.1	mg/kg
123	Arsenic	0	NA	NA	NA	NA	10.1	mg/kg
331	Arsenic	16	0.00%	5.48	8.7	1.72	10.1	mg/kg
371	Arsenic	1	0.00%	1.20	1.2	NA	10.1	mg/kg
374	Arsenic	3	0.00%	1.09	1.3	0.287	10.1	mg/kg
439	Arsenic	6	0.00%	2.05	4	1.03	10.1	mg/kg
440	Arsenic	10	10.00%	5.60	12	3.50	10.1	mg/kg
441	Arsenic	5	20.00%	7.58	11	2.23	10.1	mg/kg
442	Arsenic	11	0.00%	5.37	8.7	1.79	10.1	mg/kg
444	Arsenic	39	0.00%	4.86	8.4	1.72	10.1	mg/kg
445	Arsenic	1	0.00%	8.20	8.2	NA	10.1	mg/kg
447	Arsenic	17	0.00%	6.16	8.8	1.69	10.1	mg/kg
450	Arsenic	1	0.00%	6.30	6.3	NA	10.1	mg/kg
455	Arsenic	1	0.00%	1.70	1.7	NA	10.1	mg/kg
528	Arsenic	1	0.00%	ND	ND	ND	10.1	mg/kg
559	Arsenic	12	0.00%	4.67	8.5	1.94	10.1	mg/kg
701	Arsenic	4	0.00%	3.83	5.5	1.66	10.1	mg/kg
707	Arsenic	20	0.00%	1.49	3.3	0.614	10.1	mg/kg
712/713	Arsenic	12	33.33%	7.78	15	4.47	10.1	mg/kg
770	Arsenic	2	0.00%	5.80	6.1	0.424	10.1	mg/kg
771	Arsenic	12	25.00%	10.2	31.8	7.57	10.1	mg/kg
774	Arsenic	4	0.00%	5.53	8.2	2.16	10.1	mg/kg
776	Arsenic	20	0.00%	4.13	9	2.46	10.1	mg/kg
777	Arsenic	25	4.00%	3.47	11	2.37	10.1	mg/kg
778	Arsenic	15	0.00%	4.33	10	2.31	10.1	mg/kg
779	Arsenic	3	33.33%	7.80	11	3.67	10.1	mg/kg
865	Arsenic	25	0.00%	3.16	6.3	1.43	10.1	mg/kg
881	Arsenic	25	0.00%	4.83	7.6	1.03	10.1	mg/kg
883	Arsenic	12	0.00%	3.64	8.5	2.53	10.1	mg/kg
886	Arsenic	23	4.35%	4.20	12.3	2.36	10.1	mg/kg
889	Arsenic	3	0.00%	2.67	3.8	1.03	10.1	mg/kg
991	Arsenic	10	10.00%	4.12	12	3.81	10.1	mg/kg
X 7 4 X 7	+ Amaliaa		ND - Not Dot					

NA = Not Applicable

ND = Not Detected

¹⁰ This column indicates those samples that were detected and greater than background. As a result, if this field shows 0% and a mean and maximum concentration are reported, the values represent those analytes detected but less than background.

2.1.4 Barium

Barium was not identified or discussed in building process information (CDH 1992; DOE 2004). Barium has not been found associated with UBC sites (DOE 2004).

Barium was initially identified in the ChemRisk Task 1 Report as a chemical in inventory at RFETS (although no specific building was identified) (CDH 1991a, 1991b). Several chemical compounds were identified in the Task 1 Report, which indicated small quantities were in inventory with the exception of barium chloride, which had an inventory ranging between 9 kg (in 1988) and 23 kg (in 1974). However, based on the estimated quantity of these chemicals used, barium was not carried forward as a material of concern for the ChemRisk process.

Barium waste has been generated from both laboratory and process buildings (Table 4), and from Buildings 439 (Radiological Survey), 442 (filter test facility), 778 (Plant Laundry Facility), 886 (Critical Mass Lab), and 887 (managed process and sanitary waste) (Table 7).

Barium was not identified above a RFCA AL requiring an accelerated action based on SAPs, SAP Addenda, or Closeout Reports for IHSSs and UBC sites.

In reviewing the barium soil data beneath the slabs for UBC sites (Figure 4), the summary statistics presented in Table 11 were generated. Accelerated action closeout and data summary reports have indicated barium to be associated with building concrete.

Table 11 UBC Soil Summary Statistics for Barium (mg/kg)

						\ \ \ \ \ \ \ \ \ \ \ \ \ \ \ \ \ \ \ \		
UBC	Analyte	Total Number of Samples	Detection Frequency > Background ¹¹	Mean	Maximum Concentration	Standard Deviation	Background Mean Plus 2SD	Unit
122	Barium	6	0.00%	48.5	87	26.1	141	mg/kg
123	Barium	0	NA	NA	NA	NA	141	mg/kg
331	Barium	16	0.00%	67.7	100	17.8	141	mg/kg
371	Barium	1	0.00%	16.0	16	NA	141	mg/kg
374	Barium	3	0.00%	25.3	31.8	9.80	141	mg/kg
439	Barium	6	0.00%	52.1	78.5	14.7	141	mg/kg
440	Barium	10	10.00%	90.6	290	73.8	141	mg/kg
441	Barium	5	0.00%	71.8	82	6.98	141	mg/kg
442	Barium	11	0.00%	78.8	104	26.8	141	mg/kg
444	Barium	39	7.69%	78.0	200	35.0	141	mg/kg
445	Barium	1	0.00%	81.0	81	NA	141	mg/kg
447	Barium	17	5.88%	81.2	170	35.0	141	mg/kg
450	Barium	1	0.00%	93.0	93	NA	141	mg/kg
455	Barium	1	0.00%	52.0	52	NA	141	mg/kg
528	Barium	1	100.00%	180	180	NA	141	mg/kg
559	Barium	12	0.00%	67.0	130	32.5	141	mg/kg
701	Barium	4	0.00%	88.0	120	23.7	141	mg/kg
707	Barium	20	0.00%	45.3	91	17.4	141	mg/kg
712/713	Barium	12	8.33%	101	180	37.6	141	mg/kg

¹¹ This column indicates those samples that were detected and greater than background. As a result, if this field shows 0% and a mean and maximum concentration are reported, the values represent those analytes detected but less than background.

UBC	Analyte	Total Number of Samples	Detection Frequency > Background ¹¹	Mean	Maximum Concentration	Standard Deviation	Background Mean Plus 2SD	Unit
770	Barium	2	100.00%	175	200	35.4	141	mg/kg
771	Barium	12	16.67%	98.5	196	40.3	141	mg/kg
774	Barium	4	50.00%	145	180	31.1	141	mg/kg
776	Barium	20	0.00%	70.8	93	17.9	141	mg/kg
777	Barium	25	0.00%	72.0	140	25.1	141	mg/kg
778	Barium	15	0.00%	80.8	130	27.4	141	mg/kg
779	Barium	3	0.00%	81.7	94	18.0	141	mg/kg
865	Barium	25	12.00%	95.6	250	49.8	141	mg/kg
881	Barium	25	4.00%	73.1	232	37.1	141	mg/kg
883	Barium	12	16.67%	80.4	250	68.2	141	mg/kg
886	Barium	23	13.04%	92.0	188	38.3	141	mg/kg
889	Barium	3	0.00%	65.1	82.8	19.4	141	mg/kg
991	Barium	10	0.00%	62.6	90	14.9	141	mg/kg

NA = Not Applicable

2.1.5 Cadmium

Cadmium compounds used at RFETS include elemental or metallic cadmium oxide, cadmium chloride, and cadmium sulfate (CDH 1992).

Cadmium was used in pit construction (Building 707); however, the amounts were relatively minor compared to the primary materials used (plutonium, uranium, beryllium, aluminum, and stainless steel) (CDH 1992; DOE 2004). Cadmium was rolled and formed in Buildings 444, 883, and 865. Cadmium was also used as a plating material (as cadmium salt) for plutonium and uranium components (Buildings 776/777 and 881). Cadmium was also alloyed with other metals (Building 444). Cadmium salts were used as neutron absorbers for criticality safety in recovery operations that took place in equipment that was not dimensionally safe (Buildings 771 and 881). Cadmium was used for thermal neutron shielding.

Cadmium plating wastes were treated in Building 774. Dilute cadmium plating rinsing solutions went to Building 374. Prior to the use of Building 374, they were sent to the Solar Evaporation Ponds (SEPs) (CDH 1992)¹². Process waste containing cadmium, chromium and lead were managed in Building 731 (plenum deluge and process waste pit for Building 707) and Building 528 (process waste handling building).

Cadmium was identified in a 1988/1989 Chemical Inventory list for Buildings 123, 559 and 561 as laboratory chemicals acetate, chloride, iodide, nitrate, oxide, sulfate, and metal (CDH 1992). The Health Physics Laboratory (Building 123) identified 300 grams of cadmium metal.

¹² The maximum cadmium surface soil concentration at the SEPs was 382 mg/kg, which was below both the RFCA Tier I and II ALs (1920 mg/kg) that existed at the time of the accelerated action and below the May 2003 modifications to the RFCA ALs (962 mg/kg) (DOE 2003b). The maximum cadmium subsurface soil concentration at the SEPs was 547 mg/kg, which was below both the RFCA Tier I and II ALs (1920 mg/kg) and below the May 2003 modifications to the RFCA ALs (962 mg/kg).

All of the buildings identified above involved radiological operations and included extensive HEPA filtration systems. Any particulates or fines from machining cadmium metals would have been collected on these filters prior to release from the buildings.

Cadmium was identified in the ChemRisk reports as both a chemical (for example, cadmium nitrate) and in the elemental form and as a result was evaluated as cadmium compounds (CDH 1991a). Of the 100 kg of cadmium on the 1974 inventory, 57 percent was elemental and 34 percent was cadmium oxide (CDH 1992). Of the 46 kg of cadmium on the 1988/89 inventory, 31 percent was elemental and 56 percent was oxide.

Cadmium compounds were carried forward as materials of concern for the ChemRisk reports (CDH 1992). However, the Tasks 3 & 4 report indicated that, based on the nature of their use, cadmium compounds did not warrant further quantitative evaluation of potential off-site impacts when comparing the difference between the source maps and inventory quantities. A comparison of the emission source maps with inventory quantities presented in the building summaries (Appendix B to the Tasks 3 & 4 report) indicated buildings or processes that used cadmium were not identified as emission sources. This was due to the manner in which the material was stored, processed, or handled and was not expected to lead to significant emissions. In addition, on a number of the emission source maps, the waste treatment buildings were identified as air emission sources for chemicals that were not expected to be released in significant quantities in their primary areas of use as indicated by inventory quantities.

In addition, cadmium was one of 5 metals (cadmium, chromium, lead, mercury, and nickel) included in a group of 13 chemicals that underwent extensive investigation by ChemRisk (CDH 1992). Results indicated that uses of these materials at RFETS had been extremely limited in scope or duration, associated with insignificant quantities of the material, or involved processes or forms of the materials that were not expected to have significant off-site releases. These materials, therefore, did not warrant further quantitative evaluation as potential off-site impacts in the ChemRisk process.

Spills involving process wastes (containing cadmium, chromium, and lead) did occur within certain buildings, based on a review of RLCRs and PDSRs for these buildings. Specifically, history and process knowledge for Buildings 371, 374 and 559 revealed multiple spills of acids typically containing cadmium, chromium, and lead from the recovery of plutonium. These liquids may also have contained detectable levels of RCRA volatile organics such as carbon tetrachloride and perchloroethylene. A small number of randomly located concrete floor samples were collected and analyzed for RCRA metals and volatile organic compounds (VOCs). All results indicated concentrations were below regulatory limits.

In addition, historical process knowledge for Buildings 771 and 774 indicate toxic metals were present in solutions and sludge/residues contained in process equipment, tanks, process lines, and waste containers. There were many incidents involving nitric acid solution spills that etched into the floor or walls. However, all sample results were below RCRA regulatory limits.

Cadmium waste has been generated from both laboratory and process buildings (Table 4 and Table 7).

Cadmium was not identified above a RFCA AL requiring an accelerated action based on SAPs, SAP Addenda, or the Closeout Report for IHSSs and UBC sites.

In reviewing the cadmium soil data beneath the slabs for UBC sites (Figure 5), the summary statistics presented in Table 12 were generated.

Table 12 UBC Soil Summary Statistics for Cadmium (mg/kg)

UBC	Analyte	Total Number of Samples	Detection Frequency > Background ¹³	Mean	Maximum Concentration	Standard Deviation	Background Mean Plus 2SD	Unit
122	Cadmium	6	0.00%	ND	ND	ND	1.62	mg/kg
123	Cadmium	0	NA	NA	NA	NA	1.62	mg/kg
331	Cadmium	16	6.25%	0.169	2.2	0.542	1.62	mg/kg
371	Cadmium	1	0.00%	ND	ND	ND	1.62	mg/kg
374	Cadmium	3	0.00%	ND	ND	ND	1.62	mg/kg
439	Cadmium	6	0.00%	ND	ND	ND	1.62	mg/kg
440	Cadmium	10	0.00%	0.127	0.41	0.118	1.62	mg/kg
441	Cadmium	5	0.00%	ND	ND	ND	1.62	mg/kg
442	Cadmium	11	0.00%	0.139	0.4	0.147	1.62	mg/kg
444	Cadmium	39	5.13%	0.782	14	2.89	1.62	mg/kg
445	Cadmium	1	0.00%	0.160	0.16	NA	1.62	mg/kg
447	Cadmium	17	0.00%	0.216	1.6	0.412	1.62	mg/kg
450	Cadmium	1	0.00%	0.280	0.28	NA	1.62	mg/kg
455	Cadmium	1	0.00%	ND	ND	ND	1.62	mg/kg
528	Cadmium	1	0.00%	0.450	0.45	NA	1.62	mg/kg
559	Cadmium	12	0.00%	0.092	0.23	0.071	1.62	mg/kg
701	Cadmium	4	0.00%	ND	ND	ND	1.62	mg/kg
707	Cadmium	20	0.00%	0.055	0.27	0.061	1.62	mg/kg
712/713	Cadmium	12	8.33%	0.660	3.9	1.11	1.62	mg/kg
770	Cadmium	2	0.00%	0.308	0.58	0.384	1.62	mg/kg
771	Cadmium	12	8.33%	0.424	2.3	0.642	1.62	mg/kg
774	Cadmium	4	0.00%	0.051	0.093	0.028	1.62	mg/kg
776	Cadmium	20	0.00%	0.053	0.43	0.089	1.62	mg/kg
777	Cadmium	25	0.00%	0.095	1.1	0.229	1.62	mg/kg
778	Cadmium	15	0.00%	ND	ND	ND	1.62	mg/kg
779	Cadmium	3	0.00%	0.062	0.08	0.024	1.62	mg/kg
865	Cadmium	25	0.00%	0.045	0.17	0.034	1.62	mg/kg
881	Cadmium	25	0.00%	0.076	0.33	0.075	1.62	mg/kg
883	Cadmium	12	0.00%	0.224	1.6	0.456	1.62	mg/kg
886	Cadmium	23	0.00%	0.123	0.38	0.103	1.62	mg/kg
889	Cadmium	3	0.00%	0.075	0.11	0.045	1.62	mg/kg
991	Cadmium	10	0.00%	0.117	0.86	0.261	1.62	mg/kg

NA = Not Applicable

ND = Not Detected

¹³ This column indicates those samples that were detected and greater than background. As a result, if this field shows 0% and a mean and maximum concentration are reported, the values represent those analytes detected but less than background.

2.1.6 Chromium

Chromium compounds were used for plating in the Building 444 Research and Development (R&D) plating lab (CDH 1992; DOE 2004). Some solutions were made by mixing chromium salts with acids; others were purchased in aqueous forms. Chromium was present in anion exchange resins in Building 371. Chromium trioxide was used in Building 444 (with sulfuric and phosphoric acids) to chemically mill beryllium. Prior to 1976, chromates were added to the water as a rust inhibitor used in the Building 712/713 Cooling Towers.

Before RCRA, plating wastes were treated in Building 774 (CDH 1992). Dilute rinsates were sent to Building 374. Prior to Building 374, the Solar Evaporation Ponds were used to treat wastewater¹⁴. Process waste containing cadmium, chromium and lead were managed in Building 731 (plenum deluge and process waste pit for Building 707) and Building 528 (process waste handling building).

Chromium was identified in a 1988/1989 Chemical Inventory list for Buildings 123, 559 and 561 in various chemical forms such as chloride, nitrate, oxide, potassium sulfate, sulfate, and trioxide (CDH 1992).

Chromium compounds were carried forward as materials of concern for the ChemRisk reports (CDH 1992). However, the Tasks 3 & 4 report indicated that, based on the nature of their use, they did not warrant further quantitative evaluation of potential off-site impacts. A comparison of the emission source maps with inventory quantities presented in the building summaries (Appendix B to the Tasks 3 & 4 report) indicated buildings or processes that used chromium were not identified as emission sources. This was due to the manner in which the material was stored, processed, or handled, and was not expected to lead to significant emissions.

In addition, chromium was one of 5 metals (cadmium, chromium, lead, mercury, and nickel) included in a group of 13 chemicals that underwent extensive investigation by ChemRisk (CDH 1992). Results indicated that uses of these materials at RFETS had been extremely limited in scope or duration, associated with insignificant quantities of the material, or involved processes or forms of the materials that were not expected to have significant off-site releases. These materials, therefore, did not warrant further quantitative evaluation as potential off-site impacts in the ChemRisk process.

Spills involving process wastes (containing cadmium, chromium, and lead) did occur within certain buildings, based on a review of RLCRs and PDSRs for these buildings. Specifically, history and process knowledge for Buildings 371, 374, and 559 revealed multiple spills of acids typically containing cadmium, chromium, and lead from the recovery of plutonium. These liquids may also have contained detectable levels of RCRA volatile organics such as carbon tetrachloride and perchloroethylene. A small number of randomly located concrete floor samples were collected and analyzed for RCRA metals and VOCs. All results indicated concentrations were below regulatory limits.

¹⁴ The maximum chromium surface soil concentration at the SEPs was 120 mg/kg and the maximum subsurface soil concentration was 56.9 mg/kg (DOE 2003b). These concentrations are below both the RFCA Tier I (4.41E+05 mg/kg) and Tier II (4410 mg/kg) ALs (surface and subsurface soil) for chromium VI that existed at the time of the accelerated action. These concentrations were also below the May 2003 modifications to the RFCA ALs (268 mg/kg) for chromium VI.

444

445

447

450

Chromium (total)

Chromium (total)

Chromium (total)

Chromium (total)

In addition, historical process knowledge for Buildings 771 and 774 indicate toxic metals were present in solutions and sludge/residues contained in process equipment, tanks, process lines, and waste containers. There were many incidents involving nitric acid solution spills that etched into the floor or walls. However, all sample results were below RCRA regulatory limits.

Chromium waste has been generated from both laboratory and process buildings (Table 4 and Table 7).

Chromium was identified as present in soil above the RFCA AL requiring an accelerated action based on SAPs, SAP Addenda, or the Closeout Report for IHSS 500-158, near Building 551. (This may have been related to scrap metal storage in the area.) Immediately north of Building 551, one surface soil sampling location (CA41-025) indicated a chromium concentration (2,600 mg/kg) greater than the RFCA WRW AL of 268 mg/kg (DOE 2004c). This location was removed and all remaining concentrations were below RFCA WRW ALs (DOE 2004c).

In 1989, a chromic acid spill from the basement of Building 444 passed through the sanitary waste treatment system and reached an on-site retention pond (B-3). The solution leaked through cracks in the floor into the building foundation drain system. It was collected in a sump and pumped into the Plant's sanitary sewer system. The water was discharged to retention Pond B-3. This water was then pumped to the spray fields (IHSSs 216.2 and 216.3). Because the ponds and the surfaces of the spray fields were frozen, significant amounts of chromic acid-contaminated spray water ran off the hillsides adjacent to the spray fields. This water was collected in Ponds B-3, B-4, and B-5 on Plant site. Water from Pond B-5 was pumped into Upper Church Ditch; concentrations in this water were below the surface water standard of 0.05 part per million (ppm) (CDH 1992).

In reviewing the chromium soil data beneath the slabs for UBC sites (Figure 6), the summary statistics presented in Table 13 were generated.

UBC	Analyte	Total Number of Samples	Detection Frequency > Background ¹⁵	Mean	Maximum Concentration	Standard Deviation	Background Mean Plus 2SD	Unit
122	Chromium (total)	6	16.67%	12.2	- 17	4.89	16.8	mg/kg
123	Chromium (total)	0	NA	NA	NA	NA	16.8	mg/kg
331	Chromium (total)	16	56.25%	24.5	68	15.4	16.8	mg/kg
371	Chromium (total)	1	0.00%	6.50	6.5	NA	16.8	mg/kg
374	Chromium (total)	3	0.00%	8.70	11.8	3.10	16.8	mg/kg
439	Chromium (total)	6	0.00%	10.4	14.8	3.47	16.8	mg/kg
440	Chromium (total)	10	60.00%	18.9	37	8.52	16.8	mg/kg
441	Chromium (total)	5	60.00%	19.4	30	6.19	16.8	mg/kg
442	Chromium (total)	11	45.45%	15.9	20.7	3.73	16.8	mg/kg

18.0

27.0

19.3

14.0

79

27

27

14

11.6

NA

4.29

NA

16.8

16.8

16.8

16.8

mg/kg

mg/kg

mg/kg

mg/kg

Table 13 UBC Soil Summary Statistics for Chromium (mg/kg)

41.03%

100.00%

88.24%

0.00%

39

1

17

1

¹⁵ This column indicates those samples that were detected and greater than background. As a result, if this field shows 0% and a mean and maximum concentration are reported, the values represent those analytes detected but less than background.

UBC	Analyte	Total Number of Samples	Detection Frequency > Background 15	Mean	Maximum Concentration	Standard Deviation	Background Mean Plus 2SD	Unit
455	Chromium (total)	1	0.00%	11.0	11	NA	16.8	mg/kg
528	Chromium (total)	1	100.00%	23.0	23	NA	16.8	mg/kg
559	Chromium (total)	12	25.00%	12.7	26	5.30	16.8	mg/kg
701	Chromium (total)	4	75.00%	17.5	21	2.89	16.8	mg/kg
707	Chromium (total)	20	0.00%	9.61	15	2.18	16.8	mg/kg
712/713	Chromium (total)	12	66.67%	32.3	100	28.3	16.8	mg/kg
770	Chromium (total)	2	50.00%	22.0	29	9.90	16.8	mg/kg
771	Chromium (total)	12	25.00%	17.7	58	13.0	16.8	mg/kg
774	Chromium (total)	4	75.00%	17.8	19	1.50	16.8	mg/kg
776	Chromium (total)	20	60.00%	19.8	34	7.57	16.8	mg/kg
777	Chromium (total)	25	56.00%	29.3	160	36.0	16.8	mg/kg
778	Chromium (total)	15	33.33%	16.5	32	8.01	16.8	mg/kg
779	Chromium (total)	3	100.00%	20.0	21	1.73	16.8	mg/kg
865	Chromium (total)	25	28.00%	15.1	36	6.66	16.8	mg/kg
881	Chromium (total)	25	32.00%	18.9	111	19.5	16.8	mg/kg
883	Chromium (total)	12	41.67%	16.8	31	5.86	16.8	mg/kg
886	Chromium (total)	23	30.43%	15.0	35.2	7.76	16.8	mg/kg
889	Chromium (total)	3	66.67%	16.3	20.1	4.14	16.8	mg/kg
991	Chromium (total)	10	50.00%	16.2	32	7.75	16.8	mg/kg

NA = Not Applicable

2.1.7 Cobalt

Cobalt was not identified or discussed in building process information (CDH 1992; DOE 2004). Cobalt has not been found associated with UBC sites (DOE 2004).

Cobalt was initially identified in the ChemRisk Task 1 report as a chemical in inventory at RFETS (although no specific building was identified) (CDH 1991a). A majority of these chemicals appeared to have been used as laboratory standards or analytical testing materials because they were used in very small quantities. However, one compound, cobalt oxide was identified as present in the 1974 inventory at 677 kg and then later in 1988 in less than 1 kg.

In the ChemRisk Task 2 report (stage 2 screening) a derived reference dose (RfD) was calculated for cobalt by dividing its lethal dose (LD_{50}^{16}) by a factor of 100,000 (CDH 1991b). This approach was considered very conservative for essential nutrients like cobalt. The human daily dietary intake of cobalt was estimated to be approximately 0.1 to 0.25 milligram per day (mg/day) (based on a 1963 California State Water Resources Control Board concentration [CDH 1991b]). Using the exposure scenarios described, concentrations of cobalt in air and water were calculated. The average daily doses of cobalt received by a maximally exposed individual through inhalation and drinking water ingestion were 0.029 mg/day and 0.085 mg/day, respectively. These doses were lower than the daily dietary intake level and were unlikely to pose a health hazard to off-site individuals. For this reason, cobalt was not carried forward as a material of concern for the

¹⁶ LD₅₀ is the amount of material, given all at once, which causes the death of 50 percent (one half) of a group of test animals. LD₅₀ is one way of measuring the acute toxicity of a material.

ChemRisk process. Note that cobalt oxide was identified in the various tables during the stage 2 screening.

There is no record of spills involving cobalt within these buildings, based on a review of RLCRs and PDSRs for these buildings. There is no record of cobalt waste being generated at RFETS (Table 4 and Table 7).

Cobalt was not identified above a RFCA AL requiring an accelerated action based on SAPs, SAP Addenda, or Closeout Reports for IHSSs and UBC sites.

In reviewing the cobalt soil data beneath the slabs for UBC sites (Figure 7), the summary statistics presented in Table 14 were generated.

Table 14 UBC Soil Summary Statistics for Cobalt (mg/kg)

UBC	Analyte	Total Number of Samples	Detection Frequency > Background ¹⁷	Mean	Maximum Concentration	Standard Deviation	Background Mean Plus 2SD	Unit
122	Cobalt	6	0.00%	3.28	5.3	1.41	10.9	mg/kg
123	Cobalt	0	NA	NA	NA	NA	10.9	mg/kg
331	Cobalt	16	6.25%	6.64	22	4.45	10.9	mg/kg
371	Cobalt	1	0.00%	1.40	1.4	NA	10.9	mg/kg
374	Cobalt	3	33.33%	8.23	21.1	11.2	10.9	mg/kg
439	Cobalt	6	0.00%	4.95	9	2.18	10.9	mg/kg
440	Cobalt	10	40.00%	14.7	47	13.5	10.9	mg/kg
441	Cobalt	5	0.00%	5.64	8.1	1.50	10.9	mg/kg
442	Cobalt	11	9.09%	6.00	12.9	3.04	10.9	mg/kg
444	Cobalt	39	25.64%	11.1	61	12.0	10.9	mg/kg
445	Cobalt	1	0.00%	9.10	9.1	NA	10.9	mg/kg
447	Cobalt	17	23.53%	9.85	34	7.21	10.9	mg/kg
450	Cobalt	1	100.00%	12.0	12	NA	10.9	mg/kg
455	Cobalt	1	0.00%	5.40	5.4	NA	10.9	mg/kg
528	Cobalt	1	0.00%	2.50	2.5	NA	10.9	mg/kg
559	Cobalt	12	33.33%	9.98	26	8.09	10.9	mg/kg
701	Cobalt	4	0.00%	6.93	9.4	1.82	10.9	mg/kg
707	Cobalt	20	15.00%	8.31	36	9.19	10.9	mg/kg
712/713	Cobalt	12	16.67%	8.88	34	8.58	10.9	mg/kg
770	Cobalt	2	0.00%	7.05	8.4	1.91	10.9	mg/kg
771	Cobalt	12	8.33%	9.34	27.7	6.13	10.9	mg/kg
774	Cobalt	4	0.00%	6.90	8.4	2.30	10.9	mg/kg
776	Cobalt	20	60.00%	13.2	32	6.93	10.9	mg/kg
777	Cobalt	25	44.00%	12.1	34	7.86	10.9	mg/kg
778	Cobalt	15	40.00%	14.3	55	14.8	10.9	mg/kg
779	Cobalt	3	66.67%	8.30	11	4.68	10.9	mg/kg
865	Cobalt	25	0.00%	5.63	9.6	1.45	10.9	mg/kg
881	Cobalt	25	48.00%	18.3	137	28.6	10.9	mg/kg
883	Cobalt	12	50.00%	10.1	17	3.99	10.9	mg/kg
886	Cobalt	23	4.35%	4.75	14	2.36	10.9	mg/kg

¹⁷ This column indicates those samples that were detected and greater than background. As a result, if this field shows 0% and a mean and maximum concentration are reported, the values represent those analytes detected but less than background.

UBC	Analyte	Total Number of Samples	Detection Frequency > Background ¹⁷	Mean	Maximum Concentration	Standard Deviation	Background Mean Plus 2SD	Unit
889	Cobalt	3	33.33%	8.90	11.4	2.98	10.9	mg/kg
991	Cobalt	10	60.00%	12.2	28	6.97	10.9	mg/kg

NA = Not Applicable

2.1.8 Copper

Copper was primarily used in metallurgical operations involved in the development of alloys within Buildings 865, 881, and 883 (CDH 1992; DOE 2004). All of these buildings involved radiological operations and included extensive HEPA filtration systems. Any particulates or fines from machining copper metals would have been collected on these filters prior to release from the buildings.

Copper was identified in the ChemRisk Task 1 report as either copper cyanide or copper sulfate and not as the metal (CDH 1991a). Copper was not carried forward as a material of concern for the ChemRisk process indicating an insufficient quantity existed at RFETS to pose a potential off-site health hazard (CDH 1991b).

Copper waste was generated from Building 559 (acid containing copper) and 881 (copper cyanide) (Table 4 and Table 7).

Copper was not identified above a RFCA AL requiring an accelerated action based on Closeout Reports for IHSSs and UBC sites. However, an elevated concentration of copper greater than the RFCA action level was identified at the East Firing Range and was removed. Soil removal at the East Firing Range was based on the presence of lead and not copper.

In reviewing the copper soil data beneath the slabs for UBC sites (Figure 8), the summary statistics presented in Table 15 were generated.

Table 15 UBC Soil Summary Statistics for Copper (mg/kg)

UBC	Analyte	Total Number of Samples	Detection Frequency > Background ¹⁸	Mean	Maximum Concentration	Standard Deviation	Background Mean Plus 2SD	Unit
122	Copper	6	0.00%	9.68	14	2.97	18.1	mg/kg
123	Copper	0	NA	NA	NA	NA	18.1	mg/kg
331	Copper	16	37.50%	22.6	75	17.5	18.1	mg/kg
371	Copper	1	0.00%	4.30	4.3	NA	18.1	mg/kg
374	Copper	3	33.33%	31.5	84.1	45.6	18.1	mg/kg
439	Copper	6	33.33%	16.4	30.4	7.88	18.1	mg/kg
440	Copper	10	70.00%	40.5	120	33.8	18.1	mg/kg
441	Copper	5	0.00%	10.4	13	1.65	18.1	mg/kg
442	Copper	11	0.00%	10.7	16.7	3.99	18.1	mg/kg
444	Copper	39	48.72%	31.0	190	36.8	18.1	mg/kg

¹⁸ This column indicates those samples that were detected and greater than background. As a result, if this field shows 0% and a mean and maximum concentration are reported, the values represent those analytes detected but less than background.

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UBC	Analyte	Total Number of Samples	Detection Frequency > Background ¹⁸	Mean	Maximum Concentration	Standard Deviation	Background Mean Plus 2SD	Unit
445	Copper	1	0.00%	12.0	12	NA	18.1	mg/kg
447	Copper	17	52.94%	22.7	80	16.9	18.1	mg/kg
450	Copper	1	100.00%	25.0	25	NA	18.1	mg/kg
455	Copper	1	100.00%	41.0	41	NA	18.1	mg/kg
528	Copper	1	100.00%	60.0	60	NA	18.1	mg/kg
559	Copper	12	33.33%	24.2	67	21.3	18.1	mg/kg
701	Copper	4	75.00%	32.8	68	24.5	18.1	mg/kg
707	Copper	20	25.00%	20.8	90	22.4	18.1	mg/kg
712/713	Copper	12	91.67%	105	430	137	18.1	mg/kg
770	Copper	2	0.00%	14.5	18	4.95	18.1	mg/kg
771	Copper	12	66.67%	79.6	421	135	18.1	mg/kg
774	Copper	4	25.00%	16.0	21	3.56	18.1	mg/kg
776	Copper	20	75.00%	35.1	92	21.9	18.1	mg/kg
777	Copper	25	68.00%	31.5	93	22.7	18.1	mg/kg
778	Copper	15	46.67%	34.5	140	37.2	18.1	mg/kg
779	Copper	3	0.00%	7.60	9.8	2.31	18.1	mg/kg
865	Copper	25	8.00%	13.1	26	4.69	18.1	mg/kg
881	Copper	25	0.00%	13.3	17.9	2.79	18.1	mg/kg
883	Copper	12	83.33%	58.8	180	47.5	18.1	mg/kg
886	Copper	23	52.17%	144	1190	265	18.1	mg/kg
889	Copper	3	66.67%	47.5	73	28.2	18.1	mg/kg
991	Copper	10	70.00%	42.1	82	28.5	18.1	mg/kg

NA = Not Applicable

2.1.9 Iron

Processes involved in Buildings 444, 445, 450, and 455 included cleaning graphite crucibles, which were used for depleted uranium and beryllium metallurgy (CDH 1992). The removed residues contained trace amounts of iron and other cast metals. Uranium machining (Building 444) used parts fabricated from depleted uranium that contained trace amounts of iron, silica, titanium, aluminum, and stainless steel. Iron was associated with anion exchange resins in Building 371.

All of the buildings identified above involved radiological operations and included extensive HEPA filtration systems. Any particulates or fines from machining iron metals would have been collected on these filters prior to release from the buildings.

Cast iron was also the material used in pipe construction for various process waste lines within the former IA.

Iron as a metal was not identified in the ChemRisk Task 1 report as a material in inventory at RFETS (CDH 1991a). Various iron chemical compounds were identified such as ferrous sulfide, ferric nitrate, ferrous ammonium sulfate and ferrous sulfamate. Quantities ranged between 0.5 to approximately 450 kg in the 1974 inventory and 5 to 900 kg in 1988 inventory. However, iron was not carried forward as a material of concern for the ChemRisk process (CDH 1991b).

There is no record of iron waste being generated at RFETS (Table 4 and Table 7).

Iron was not identified above a RFCA AL requiring an accelerated action based on SAPs, SAP Addenda, or Closeout Reports for IHSSs and UBC sites.

In reviewing the iron soil data beneath the slabs for UBC sites (Figure 9), the summary statistics presented in Table 16 were generated.

Table 16 UBC Soil Summary Statistics for Iron (mg/kg)

	Table 10 OBC Soil Summary Statistics for fron (mg/kg)								
UBC	Analyte	Total Number of Samples	Detection Frequency > Background ¹⁹	Mean	Maximum Concentration	Standard Deviation	Background Mean Plus 2SD	Unit	
122	Iron	6	0.00%	11,367	15000	3,837	17,601	mg/kg	
123	Iron	0	NA	NA	NA	NA	17,601	mg/kg	
331	Iron	16	12.50%	14,956	21000	2,998	17,601	mg/kg	
371	Iron	1	0.00%	7,200	7200	NA	17,601	mg/kg	
374	Iron	3	0.00%	10,367	17300	6,123	17,601	mg/kg	
439	Iron	6	16.67%	12,517	24300	5,960	17,601	mg/kg	
440	Iron	10	10.00%	12,500	18000	3,624	17,601	mg/kg	
441	Iron	5	20.00%	15,800	19000	2,387	17,601	mg/kg	
442	Iron	11	9.09%	13,415	17900	3,246	17,601	mg/kg	
444	Iron	39	12.82%	14,310	26000	4,243	17,601	mg/kg	
445	Iron	1	100.00%	21,000	21000	NA	17,601	mg/kg	
447	Iron	17	52.94%	18,647	29000	4,609	17,601	mg/kg	
450	Iron	1	0.00%	12,000	12000	NA	17,601	mg/kg	
455	Iron	1	0.00%	11,000	11000	NA	17,601	mg/kg	
528	Iron	1	0.00%	5,400	5400	NA	17,601	mg/kg	
559	Iron	12	8.33%	11,192	19000	3,853	17,601	mg/kg	
701	Iron	4	25.00%	15,500	21000	3,697	17,601	mg/kg	
707	Iron	20	0.00%	9,865	13000	1,800	17,601	mg/kg	
712/713	Iron	12	58.33%	20,567	54900	12,287	17,601	mg/kg	
770	Iron	2	0.00%	14,500	16000	2,121	17,601	mg/kg	
771	Iron	12	16.67%	15,000	19200	2,417	17,601	mg/kg	
774	Iron	4	0.00%	14,000	16000	1,414	17,601	mg/kg	
776	Iron	20	15.00%	13,015	19000	3,250	17,601	mg/kg	
777	Iron	25	8.00%	13,048	22000	3,513	17,601	mg/kg	
778	Iron	15	13.33%	12,473	21000	4,899	17,601	mg/kg	
779	Iron	3	100.00%	19,333	20000	1,155	17,601	mg/kg	
865	Iron	25	16.00%	13,564	24000	3,726	17,601	mg/kg	
881	Iron	25	8.00%	13,340	18300	2,444	17,601	mg/kg	
883	Iron	12	66.67%	21,242	30000	8,255	17,601	mg/kg	
886	Iron	23	8.70%	11,881	33000	5,567	17,601	mg/kg	
889	Iron	3	66.67%	22,867	31300	8,712	17,601	mg/kg	
991	Iron	10	30.00%	14,510	25000	4,731	17,601	mg/kg	

NA = Not Applicable

¹⁹ This column indicates those samples that were detected and greater than background. As a result, if this field shows 0% and a mean and maximum concentration are reported, the values represent those analytes detected but less than background.

2.1.10 Lead

Lead was mainly used for radiation shielding for plutonium operations (Building 300, 559 and 700) (CDH 1992; DOE 2004). A 1974 inventory indicated over 1 million pounds of lead. Molten lead was identified in Building 865. Lead was used for non-destructive testing in Building 460. Lead fluoride and lead metal were used in Building 771 for laboratory-scale attempts at lead/americium alloying. Lead fluoroborate and lead oxide were used in small quantities in plating operations. Process waste containing cadmium, chromium and lead were managed in Building 731 (plenum deluge and process waste pit for Building 707) and Building 528 (process waste handling building). Lead was also discharged as bullets at the East and North Firing Ranges. Lead gaskets were used in some of the older pipelines, mainly process waste and sanitary sewer lines.

Metallic lead was not considered a source of contamination in the ChemRisk reports. In the ChemRisk Task 1 report, several lead compounds were identified in a 1988/1989 chemical inventory list for Buildings 123, 559, 561, and 701 including acetate, chloride, iodide, metal, nitrate, oxide, and powder (CDH 1992). Lead compounds were carried forward as materials of concern for the ChemRisk reports. However, the Tasks 3 & 4 report indicated that based on the nature of their use they did not warrant further quantitative evaluation of potential off-site impacts. A comparison of the emission source maps with inventory quantities presented in the building summaries (Appendix B to the Tasks 3 & 4 report) indicated buildings or processes that used lead were not identified as emission sources. This was due to the manner in which the material was stored, processed, or handled and was not expected to lead to significant emissions.

In addition, lead was one of 5 metals (cadmium, chromium, lead, mercury, and nickel) included in a group of 13 chemicals that underwent extensive investigation by ChemRisk (CDH 1992). Results indicated that uses of these materials at RFETS had been extremely limited in scope or duration, associated with insignificant quantities of the material, or involved processes or forms of the materials, which were not expected to have significant off-site releases. These materials, therefore, did not warrant further quantitative evaluation as potential off-site impacts in the ChemRisk process.

Spills involving process wastes (containing cadmium, chromium, and lead) did occur within certain buildings, based on a review of RLCRs and PDSRs for these buildings. Specifically, history and process knowledge for Buildings 371, 374, and 559 revealed multiple spills of acids typically containing cadmium, chromium, and lead from the recovery of plutonium. These liquids may also have contained detectable levels of RCRA volatile organics such as carbon tetrachloride and perchloroethylene. A small number of randomly located concrete floor samples were collected and analyzed for RCRA metals and VOCs. All results indicated concentrations were below regulatory limits.

In addition, historical process knowledge for Buildings 771 and 774 indicate toxic metals were present in solutions and sludge/residues contained in process equipment, tanks, process lines, and waste containers. There were many incidents involving nitric acid solution spills that etched into the floor or walls. However, all sample results were below RCRA regulatory limits.

Lead waste has been generated from both laboratory and process buildings (Table 4 and Table 7).

Lead was identified as present in soil above the RFCA AL requiring an accelerated action based on SAPs, SAP Addenda, or Closeout Reports (and ER RFCA Standard Operating Protocol [RSOP] notifications for routine soil remediation) for IHSSs and UBC sites; specifically, lead was identified in soil at an underground storage tank associated with Building 441 (IHSS 400-128); UBC 123, and at both the East Firing Range (IHSS SE-1602) and North Firing Range (IHSS NW-1505). The lead concentrations at UBC 123 were associated with a lead-lined sump (not from a Site process), and the lead concentrations at the East and North Firing Ranges was due to the presence of discharged lead bullets (also not associated with a Site process).

In reviewing the lead soil data beneath the slabs for UBC sites (Figure 10), the summary statistics presented in Table 17 were generated.

Table 17 UBC Soil Summary Statistics for Lead (mg/kg)

		Total	Detection	•		G: 1 1	Background	
UBC	Analyte	Number of	Frequency	Mean	Maximum	Standard Deviation	Mean Plus	Unit
		Samples	> Background ²⁰		Concentration		2SD	
122	Lead	6	0.00%	12.3	34	11.3	54.6	mg/kg
123	Lead	0	NA	NA	NA	NA	54.6	mg/kg
331	Lead	16	12.50%	25.4	170	43.4	54.6	mg/kg
371	Lead	1	0.00%	3.40	3.4	NA	54.6	mg/kg
374	Lead	3	0.00%	4.53	5.6	1.16	54.6	mg/kg
439	Lead	6	0.00%	13.1	24	7.51	54.6	mg/kg
440	Lead	10	0.00%	9.09	12	2.26	54.6	mg/kg
441	Lead	5	20.00%	18.9	55	20.2	54.6	mg/kg
442	Lead	11	9.09%	16.8	56.3	17.0	54.6	mg/kg
444	Lead	39	2.56%	49.2	1500	239	54.6	mg/kg
445	Lead	1	0.00%	12.0	12	NA	54.6	mg/kg
447	Lead	17	17.65%	78.0	590	160	54.6	mg/kg
450	Lead	1	0.00%	8.20	8.2	NA	54.6	mg/kg
455	Lead	Ī .	0.00%	13.0	13	NA	54.6	mg/kg
528	Lead	1	0.00%	25.0	25	NA	54.6	mg/kg
559	Lead	12	0.00%	8.93	17	3.66	54.6	mg/kg
701	Lead	4	0.00%	15.3	21	4.19	54.6	mg/kg
707	Lead	20	0.00%	7.75	15	2.21	54.6	mg/kg
712/713	Lead	12	8.33%	21.9	69	19.2	54.6	mg/kg
770	Lead	2	0.00%	15.0	17	2.83	54.6	mg/kg
771	Lead	12	0.00%	14.5	22.4	5.24	54.6	mg/kg
774	Lead	4	0.00%	16.0	19	2.58	54.6	mg/kg
776	Lead	20	5.00%	13.2	79	15.8	54.6	mg/kg
777	Lead	25	4.00%	25.7	330	64.5	54.6	mg/kg
778	Lead	15	0.00%	8.33	14	2.36	54.6	mg/kg
779	Lead	3	0.00%	8.13	8.8	0.702	54.6	mg/kg
865	Lead	25	4.00%	20.2	250	48.1	54.6	mg/kg
881	Lead	25	0.00%	8.94	16.3	3.73	54.6	mg/kg
883	Lead	12	0.00%	13.5	34	7.75	54.6	mg/kg
886	Lead	23	0.00%	7.95	27.9	4.91	54.6	mg/kg
889	Lead	3	0.00%	9.97	11.6	1.70	54.6	mg/kg

²⁰ This column indicates those samples that were detected and greater than background. As a result, if this field shows 0% and a mean and maximum concentration are reported, the values represent those analytes detected but less than background.

374

439

Lithium

Lithium

3

6

UBC	Analyte	Total Number of Samples	Detection Frequency > Background ²⁰	Mean	Maximum Concentration	Standard Deviation	Background Mean Plus 2SD	Unit
991	Lead	10	0.00%	11.4	26	6.31	54.6	mg/kg

NA = Not Applicable

2.1.11 Lithium

Lithium was not identified or discussed in building process information in the ChemRisk Task 3 & 4 report (CDH 1992). Lithium was identified as associated with mass spectrometry analysis performed in Building 559 and as being used in Building 881 for metalworking (DOE 2004). There are a few lithium sites such as at the 903 Pad area (IHSS 140), an area outside Building 331 (IHSS 134S), and IHSS Group 300-1.

Lithium was identified in the ChemRisk Task 1 report as various chemical compounds including lithium metal (CDH 1991a). Lithium was not carried forward as a material of concern for the ChemRisk reports because it was unlikely it would have posed a reproductive hazard to off-site individuals based on the quantity of the material used (CDH 1991b).

There is no record of spills involving lithium within these buildings, based on a review of RLCRs and PDSRs for these buildings.

Lithium waste has been generated from both laboratory and process buildings (Table 4 and Table 7). The waste for a majority of the buildings (6 out of 9) was lithium batteries. The remaining three buildings (559, 881 and 883) generated a processing agent that may have contained lithium aluminum hydride, lithium carbonate and alkaline lithium metal.

Lithium metal was treated in on-site disposal areas (IHSS 140-Reative Metals Destruction Site and IHSS 134-Lithium Metal Destruction Site), however lithium was not identified above a RFCA AL requiring an accelerated action at these IHSSs and based on SAPs, SAP Addenda, or Closeout Reports for additional IHSSs and UBC sites.

In reviewing the lithium soil beneath the slabs for UBC sites (Figure 11), the summary statistics presented in Table 18 were generated.

Total Detection Background Maximum Standard **UBC** Analyte Number of Frequency Mean Mean Plus Unit Concentration Deviation > Background²¹ Samples 2SD 122 Lithium 16.67% 7.83 12 6 3.50 11.4 mg/kg 123 Lithium 0 NA NA NA NA 11.4 mg/kg 331 Lithium 16 0.00% 8.26 11 1.33 11.4 mg/kg 371 Lithium 1 0.00% 3.70 3.7 NA 11.4 mg/kg

4.83

7.17

6.6

9.6

2.04

1.89

11.4

11.4

mg/kg

mg/kg

Table 18 UBC Soil Summary Statistics for Lithium (mg/kg)

0.00%

0.00%

²¹ This column indicates those samples that were detected and greater than background. As a result, if this field shows 0% and a mean and maximum concentration are reported, the values represent those analytes detected but less than background.

UBC	Analyte	Total Number of Samples	Detection Frequency > Background ²¹	Mean	Maximum Concentration	Standard Deviation	Background Mean Plus 2SD	Unit
440	Lithium	10	40.00%	10.1	15	3.69	11.4	mg/kg
441	Lithium	5	100.00%	13.8	16	1.79	11.4	mg/kg
442	Lithium	11	45.45%	11.0	16.9	3.52	11.4	mg/kg
444	Lithium	39	43.59%	10.7	17	3.18	11.4	mg/kg
445	Lithium	1	100.00%	16.0	16	NA	11.4	mg/kg
447	Lithium	17	70.59%	12.9	20	3.11	11.4	mg/kg
450	Lithium	1	100.00%	13.0	13	NA	11.4	mg/kg
455	Lithium	1	0.00%	8.00	8	NA	11.4	mg/kg
528	Lithium	1	0.00%	10.0	10	NA	11.4	mg/kg
559	Lithium	12	8.33%	7.88	13	2.89	11.4	mg/kg
701	Lithium	4	0.00%	9.75	11	1.21	11.4	mg/kg
707	Lithium	20	5.00%	6.19	13	2.30	11.4	mg/kg
712/713	Lithium	12	41.67%	11.6	22	5.33	11.4	mg/kg
770	Lithium	2	100.00%	14.0	16	2.83	11.4	mg/kg
771	Lithium	12	16.67%	9.10	17	3.29	11.4	mg/kg
774	Lithium	4	50.00%	13.0	17	3.65	11.4	mg/kg
776	Lithium	20	40.00%	11.3	18	3.36	11.4	mg/kg
777	Lithium	25	28.00%	10.2	23	3.62	11.4	mg/kg
778	Lithium	15	33.33%	9.42	18	4.98	11.4	mg/kg
779	Lithium	3	100.00%	23.0	32	8.54	11.4	mg/kg
865	Lithium	25	20.00%	10.3	21	3.97	11.4	mg/kg
881	Lithium	25	48.00%	11.6	15.1	1.58	11.4	mg/kg
883	Lithium	12	58.33%	12.4	18	3.71	11.4	mg/kg
886	Lithium	23	21.74%	8.44	14	3.18	11.4	mg/kg
889	Lithium	3	33.33%	10.8	11.5	0.751	11.4	mg/kg
991	Lithium	10	40.00%	11.0	20	4.29	11.4	mg/kg

NA = Not Applicable

2.1.12 Manganese

Manganese was not identified or discussed in building process information (CDH 1992; DOE 2004). Manganese has not been found associated with UBC sites (DOE 2004).

Manganese was initially identified in the ChemRisk Task 1 report as a chemical in inventory at RFETS (although no specific building was identified). Several manganese chemical compounds were identified in the Task 1 report in the form of carbonate, chips, dioxide, flake, II oxide, metal, monoxide, powder, chloride, nitrate and sulfate (CDH 1991a, 1991b). Only small quantities were identified to be in inventory, with the exception of manganous sulfate which had an inventory in 1974 of 2,560 kg, and then later in 1988 of 0.06 kg. It is assumed that manganese was carried forward in the ChemRisk process based on the quantity of sulfate.

In the ChemRisk Task 2 report (stage 2 screening), manganese was identified as an essential nutrient for humans. The safe and adequate dietary allowance of manganese recommended for an adult was 10 mg/day (Based on a 1963 California State Water Resources Control Board concentration [CDH 1991b]). Using the exposure scenarios described, concentrations of manganese in air and water were calculated. Based on these doses, it was determined that

manganese was unlikely to pose a health hazard to off-site individuals and was not carried forward as a material of concern for the ChemRisk process. In addition, the predicted air concentration was approximately 1,000 times lower than the occupational air standard.

There is no record of manganese waste being generated at RFETS (Table 4 and Table 7).

Manganese was not identified above a RFCA AL requiring an accelerated action based on Closeout Reports for IHSSs and UBC sites.

In reviewing the manganese soil data beneath the slabs for UBC sites (Figure 12), the summary statistics presented in Table 19 were generated.

Table 19 UBC Soil Summary Statistics for Manganese (mg/kg)

Table 19 ODC Sulfilliary Statistics for Manganese (mg/kg)											
UBC	Analyte	Total Number of Samples	Detection Frequency > Background ²²	Mean	Maximum Concentration	Standard Deviation	Background Mean Plus 2SD	Unit			
122	Manganese	6	0.00%	108	170	36.8	365	mg/kg			
123	Manganese	0	NA	NA	NA	NA	365	mg/kg			
331	Manganese	16	0.00%	125	190	30.8	365	mg/kg			
371	Manganese	1	0.00%	63.0	63	NA	365	mg/kg			
374	Manganese	3	0.00%	156	333	155	365	mg/kg			
439	Manganese	6	16.67%	224	484	132	365	mg/kg			
440	Manganese	10	0.00%	159	250	55.3	365	mg/kg			
441	Manganese	5	0.00%	134	180	30.5	365	mg/kg			
442	Manganese	11	0.00%	147	247	71.8	365	mg/kg			
444	Manganese	39	7.69%	216	670	113	365	mg/kg			
445	Manganese	1	0.00%	100	100	NA	365	mg/kg			
447	Manganese	17	5.88%	216	550	103	365	mg/kg			
450	Manganese	1	0.00%	180	180	NA	365	mg/kg			
455	Manganese	1	0.00%	190	190	NA	365	mg/kg			
528	Manganese	1	0.00%	15.0	15	NA	365	mg/kg			
559	Manganese	12	8.33%	166	430	102	365	mg/kg			
701	Manganese	4	0.00%	195	240	31.1	365	mg/kg			
707	Manganese	20	0.00%	133	310	55.1	365	mg/kg			
712/713	Manganese	12	8.33%	211	417	87.1	365	mg/kg			
770	Manganese	2	0.00%	255	280	35.4	365	mg/kg			
771	Manganese	12	0.00%	150	229	51.7	365	mg/kg			
774	Manganese	4	0.00%	166	230	97.2	365	mg/kg			
776	Manganese	20	0.00%	177	270	47.5	365	mg/kg			
777	Manganese	25	8.00%	204	550	93.9	365	mg/kg			
778	Manganese	15	13.33%	226	660	150	365	mg/kg			
779	Manganese	3	0.00%	179	270	109	365	mg/kg			
865	Manganese	25	4.00%	213	510	86.9	365	mg/kg			
881	Manganese	25	0.00%	176	276	41.7	365	mg/kg			
883	Manganese	12	58.33%	392	650	233	365	mg/kg			
886	Manganese	23	0.00%	160	317	66.0	365	mg/kg			
889	Manganese	3	66.67%	425	556	172	365	mg/kg			

²² This column indicates those samples that were detected and greater than background. As a result, if this field shows 0% and a mean and maximum concentration are reported, the values represent those analytes detected but less than background.

UBC	Analyte	Total Number of Samples	Detection Frequency > Background ²²	Mean	Maximum Concentration	Standard Deviation	Background Mean Plus 2SD	Unit
991	Manganese	10	10.00%	176	420	92.2	365	mg/kg

NA = Not Applicable

2.1.13 Mercury

Mercury used at RFETS was, for the most part, limited to the metallic mercury contained in instruments such as barometers, manometers, and thermometers; plant machinery; mercury switches; and experimental apparatus (CDH 1992). Mercury was not used in production processes. A welding operation in Building 777 used mercury to make contact with spinning parts during welding. Mercury was collected from Plant sources and purified by distillation at Building 881; the General Laboratory. It was recycled back to the originating area in 5-pound containers.

Mercury, with the following exceptions, has not been found associated with UBC sites (DOE 2004). Mercury was found at Building 443 in the subfloor piping. It did not occur outside the pipes. Its presence was expected to be as a result from a broken gauge. There was also a report of a broken mercury gauge in Building 447 (steam plant).

Mercury was identified in the ChemRisk reports as both a chemical and metal. Materials present in 1971 included mercuric chloride, mercuric oxide, mercury/thallium batteries, electrodes, fluorescent lamps, and rectifiers (CDH 1991a). Mercury identified in a 1988/1989 chemical inventory list for Buildings 559, 561, and 886 included acetate, chloride, iodide, nitrate, oxide, sulfate and metal.

Mercury compounds were carried forward as materials of concern for the ChemRisk reports (CDH 1992). However, the Tasks 3 & 4 report indicated that, based on the nature of their use, they did not warrant further quantitative evaluation of potential off-site impacts. A comparison of the emission source maps with inventory quantities presented in the building summaries (Appendix B to the Tasks 3 & 4 report) indicated buildings or processes that used mercury were not identified as emission sources. This was due to the manner in which the material was stored, processed, or handled and was not expected to lead to significant emissions. In addition, on a number of the emission source maps, the waste treatment buildings were identified as air emission sources for chemicals that were not expected to be released in significant quantities in their primary areas of use as indicated by inventory.

In addition, mercury was one of 5 metals (cadmium, chromium lead, mercury, and nickel) included in a group of 13 chemicals that underwent extensive investigation by ChemRisk (CDH 1992). Results indicated uses of these materials at RFETS had been extremely limited in scope or duration, associated with insignificant quantities of the material, or involved processes or forms of the materials that were not expected to have significant off-site releases. These materials, therefore, did not warrant further quantitative evaluation as potential off-site impacts in the ChemRisk process.

There is no record of spills involving mercury compounds within a majority of these buildings, based on a review of RLCRs and PDSRs for these buildings. Mercury was identified as a spill

within Building 774; however, mercury was not identified as a contaminant of concern for this building because it was expected that this spill was properly remediated (K-H 1998).

Mercury waste has been generated from both laboratory and process buildings (Table 4 and Table 7).

Mercury was not identified above a RFCA AL requiring an accelerated action based on SAPs, SAP Addenda, or Closeout Reports for IHSSs and UBC sites.

In reviewing the mercury soil data beneath the slabs for UBC sites (Figure 13), the summary statistics presented in Table 20 were generated.

Table 20 UBC Soil Summary Statistics for Mercury (mg/kg)

Total Detection Standard Background											
UBC	Analyte	Number of	Frequency	Mean	Maximum	Standard	Mean Plus	Unit			
OBC	Allaryte	Samples	> Background ²³	1410411	Concentration	Deviation	2SD				
122	Mercury	6	0.00%	0.036	0.1	0.039	0.133	mg/kg			
123	Mercury	0	NA	NA	NA	NA	0.133	mg/kg			
331	Mercury	16	0.00%	0.053	0.11	0.026	0.133	mg/kg			
371	Mercury	0	NA	NA	NA	NA	0.133	mg/kg			
374	Mercury	1	0.00%	ND	ND	ND	0.133	mg/kg			
439	Mercury	6	0.00%	0.009	0.0088	0.008	0.133	mg/kg			
440	Mercury	10	0.00%	0.027	0.065	0.022	0.133	mg/kg			
441	Mercury	5	80.00%	0.330	0.58	0.226	0.133	mg/kg			
442	Mercury	11	0.00%	0.043	0.13	0.040	0.133	mg/kg			
444	Mercury	39	2.56%	0.033	0.22	0.040	0.133	mg/kg			
445	Mercury	1	0.00%	0.095	0.095	NA	0.133	mg/kg			
447	Mercury	17	0.00%	0.038	0.081	0.021	0.133	mg/kg			
450	Mercury	1	0.00%	0.046	0.046	NA	0.133	mg/kg			
455	Mercury	1	0.00%	0.009	0.0091	NA	0.133	mg/kg			
528	Mercury	1	0.00%	0.081	0.081	NA	0.133	mg/kg			
559	Mercury	12	0.00%	0.051	0.1	0.026	0.133	mg/kg			
701	Mercury	4	0.00%	0.050	0.084	0.029	0.133	mg/kg			
707	Mercury	20	0.00%	0.011	0.069	0.015	0.133	mg/kg			
712/713	Mercury	12	33.33%	0.091	0.22	0.072	0.133	mg/kg			
770	Mercury	2	0.00%	0.021	0.028	0.010	0.133	mg/kg			
771	Mercury	12	0.00%	0.039	0.082	0.023	0.133	mg/kg			
774	Mercury	4	0.00%	0.053	0.096	0.030	0.133	mg/kg			
776	Mercury	20	0.00%	0.030	0.098	0.030	0.133	mg/kg			
777	Mercury	25	4.00%	0.026	0.18	0.041	0.133	mg/kg			
778	Mercury	15	0.00%	0.015	0.029	0.009	0.133	mg/kg			
779	Mercury	3	66.67%	0.137	0.15	0.015	0.133	mg/kg			
865	Mercury	24	0.00%	0.011	0.028	0.007	0.133	mg/kg			
881	Mercury	25	0.00%	0.017	0.087	0.020	0.133	mg/kg			
883	Mercury	12	8.33%	0.031	0.14	0.039	0.133	mg/kg			
886	Mercury	23	4.35%	0.026	0.15	0.028	0.133	mg/kg			
889	Mercury	3	0.00%	0.008	0.016	0.008	0.133	mg/kg			

²³ This column indicates those samples that were detected and greater than background. As a result, if this field shows 0% and a mean and maximum concentration are reported, the values represent those analytes detected but less than background.

UBC	Analyte	Total Number of Samples	Detection Frequency > Background ²³	Mean	Maximum Concentration	Standard Deviation	Background Mean Plus 2SD	Unit
991	Mercury	8	0.00%	0.040	0.11	0.041	0.133	mg/kg

NA = Not Applicable

ND = Not Detected

2.1.14 Molybdenum

The Zero Power Plutonium Reactor (ZPPR or "zipper") project manufactured stainless steel clad fuel elements consisting of plutonium, molybdenum, and uranium from 1967 to 1968 (CDH 1992). The ZPPR fuel elements were made first by alloying the uranium and molybdenum in Building 444. The alloy was then sent to Building 771, where it was alloyed with Plutonium by casting into plates of various sizes. The ternary alloy plates were clad in stainless steel envelopes in Buildings 776/777 and sealed by welding.

The metallurgical operations in Building 865 began in 1970 and involved the development of alloys (CDH 1992). Some of the metals employed in the alloying development included aluminum, copper, magnesium, molybdenum, niobium, platinum, stainless steel, tantalum, titanium, and vanadium.

The buildings identified above involved radiological operations and included extensive HEPA filtration systems. Any emissions from machining molybdenum would have been collected on these filters prior to release from the buildings.

Molybdenum was initially identified in the ChemRisk Task 1 report as various chemical compounds in inventory at RFETS (CDH 1991b). Based on the estimated quantity of these chemicals used, molybdenum was not carried forward as a material of concern for the ChemRisk process.

Acid waste containing molybdenum was generated from Building 559.

Molybdenum was not identified above a RFCA AL requiring an accelerated action based on SAPs, SAP Addenda, or Closeout Reports for IHSSs and UBC sites.

In reviewing the molybdenum soil data beneath the slabs for UBC sites (Figure 14), the summary statistics presented in Table 21 were generated.

Table 21 UBC Soil Summary Statistics for Molybdenum (mg/kg)

UBC	Analyte	Total Number of Samples	Detection Frequency > Background ²⁴	Mean	Maximum Concentration	Standard Deviation	Background Mean Plus 2SD	Unit
122	Molybdenum	6	16.67%	0.677	1	0.237	0.941	mg/kg
123	Molybdenum	0	NA	NA	NA	NA	0.941	mg/kg
331	Molybdenum	16	18.75%	0.766	1.2	0.244	0.941	mg/kg
371	Molybdenum	1	0.00%	0.190	0.19	NA	0.941	mg/kg
374	Molybdenum	3	0.00%	0.500	0.41	0.363	0.941	mg/kg

²⁴ This column indicates those samples that were detected and greater than background. As a result, if this field shows 0% and a mean and maximum concentration are reported, the values represent those analytes detected but less than background.

UBC	Analyte	Total Number of Samples	Detection Frequency > Background ²⁴	Mean	Maximum Concentration	Standard Deviation	Background Mean Plus 2SD	Unit
439	Molybdenum	6	0.00%	0.373	0.66	0.209	0.941	mg/kg
440	Molybdenum	10	0.00%	0.540	0.84	0.220	0.941	mg/kg
441	Molybdenum	5	0.00%	0.370	0.55	0.131	0.941	mg/kg
442	Molybdenum	11	9.09%	0.266	1.3	0.350	0.941	mg/kg
444	Molybdenum	39	5.13%	0.375	1	0.234	0.941	mg/kg
445	Molybdenum	1	0.00%	0.690	0.69	NA	0.941	mg/kg
447	Molybdenum	17	11.76%	0.472	0.99	0.301	0.941	mg/kg
450	Molybdenum	1	0.00%	ND	ND	ND	0.941	mg/kg
455	Molybdenum	1	0.00%	ND	ND	ND	0.941	mg/kg
528	Molybdenum	1	0.00%	ND	ND	ND	0.941	mg/kg
559	Molybdenum	12	8.33%	0.485	0.96	0.264	0.941	mg/kg
701	Molybdenum	4	0.00%	0.413	0.69	0.293	0.941	mg/kg
707	Molybdenum	20	15.00%	0.715	2.9	0.559	0.941	mg/kg
712/713	Molybdenum	12	25.00%	1.41	9.7	2.63	0.941	mg/kg
770	Molybdenum	2	0.00%	0.580	0.72	0.198	0.941	mg/kg
771	Molybdenum	12	25.00%	0.593	1.6	0.538	0.941	mg/kg
774	Molybdenum	4	0.00%	0.380	0.86	0.349	0.941	mg/kg
776	Molybdenum	20	10.00%	0.414	1.5	0.385	0.941	mg/kg
777	Molybdenum	25	12.00%	0.847	9.5	1.86	0.941	mg/kg
778	Molybdenum	15	6.67%	0.349	1.2	0.299	0.941	mg/kg
779	Molybdenum	3	0.00%	0.553	0.83	0.246	0.941	mg/kg
865	Molybdenum	25	4.00%	0.251	1	0.211	0.941	mg/kg
881	Molybdenum	25	4.00%	0.197	1.5	0.309	0.941	mg/kg
883	Molybdenum	12	50.00%	1.01	2.4	0.524	0.941	mg/kg
886	Molybdenum	23	8.70%	0.492	3.9	0.794	0.941	mg/kg
889	Molybdenum	3	0.00%	0.180	0.39	0.182	0.941	mg/kg
991	Molybdenum	10	0.00%	0.324	0.9	0.252	0.941	mg/kg

NA = Not Applicable

ND = Not Detected

2.1.15 Nickel

Nickel carbonyl plating was conducted in Buildings 771, 777, and 779 from the early 1950s until the early 1960s or 1970s (CDH 1992). Nickel plating by nickel carbonyl decomposition was used for uranium and delta phase (alloyed) plutonium. The waste chemistry group (Building 881 R&D) supported the Joining Technology Department to join non-nuclear metals including beryllium and in some cases using brazing alloys including nickel. Nickel plating of weapon components was conducted in Building 444 up until shutdown of the plating lab in 1990. Some plating solutions were made by mixing metal salts with acids, others were purchased in aqueous form. Nickel plating solutions were heated and used in 75-gallon tanks. Some liquid evaporated; however, measurements showed that the metals did not.

Before RCRA, plating wastes were treated in Building 774 (CDH 1992). Dilute rinsates were sent to Building 374. Prior to Building 374, the Solar Evaporation Ponds were used to treat wastewater. Building 991 was used as the product warehouse where components containing plutonium, uranium, and nickel were assembled into final products.

Nickel was found to be associated with anion exchange resins in Building 371 (DOE 2004).

The buildings identified above involved radiological operations and included extensive HEPA filtration systems. Any emissions from machining or plating nickel would have been collected on these filters prior to release from the buildings.

Nickel compounds were carried forward as materials of concern for the ChemRisk reports (CDH 1992). However, the Tasks 3 & 4 report indicated that, based on the nature of their use, they did not warrant further quantitative evaluation of potential off-site impacts. A comparison of the emission source maps with inventory quantities presented in the building summaries (Appendix B to the Tasks 3 & 4 report) indicated buildings or processes that used nickel were not identified as emission sources. This was due to the manner in which the material was stored, processed, or handled and was not expected to lead to significant emissions. In addition, on a number of the emission source maps, the waste treatment buildings were identified as air emission sources for chemicals that were not expected to be released in significant quantities in their primary areas of use as indicated by the inventory.

In addition, nickel was one of 5 metals (cadmium, chromium lead, mercury, and nickel) included in a group of 13 chemicals that underwent extensive investigation by ChemRisk (CDH 1992). Results indicated uses of these materials at RFETS had been extremely limited in scope or duration, associated with insignificant quantities of the material, or involved processes or forms of the materials that were not expected to have significant off-site releases. These materials therefore did not warrant further quantitative evaluation as potential off-site impacts in the ChemRisk process.

There is no record of spills involving nickel compounds within these buildings, based on a review of RLCRs and PDSRs for these buildings.

Waste containing nickel as an underlying hazardous constituent has been generated from both laboratory and process buildings (Table 4 and Table 7).

Nickel carbonyl canisters were stored and/or vented outside at three locations. Nickel was not identified above a RFCA AL requiring an accelerated action based on SAPs, SAP Addenda, or Closeout Reports for IHSSs and UBC sites.

In reviewing the nickel soil data beneath the slabs for UBC sites (Figure 15), the summary statistics presented in Table 22 were generated.

					•		\ \ \ \ \ \ \ \ \ \ \ \ \ \ \ \ \ \ \		
	UBC	Analyte	Total Number of Samples	Detection Frequency > Background ²⁵	Mean	Maximum Concentration	Standard Deviation	Background Mean Plus 2SD	Unit
	122	Nickel	6	16.67%	9.08	16	5.29	14.8	mg/kg
	123	Nickel	0	NA	NA	NA	NA	14.8	mg/kg
	331	Nickel	16	50.00%	16.4	38	7.79	14.8	mg/kg
	371	Nickel	1	0.00%	2.80	2.8	NA	14.8	mg/kg
Ī	374	Nickel	3	33.33%	7.57	15.7	7.13	14.8	mg/kg

Table 22 UBC Soil Summary Statistics for Nickel (mg/kg)

²⁵ This column indicates those samples that were detected and greater than background. As a result, if this field shows 0% and a mean and maximum concentration are reported, the values represent those analytes detected but less than background.

UBC	Analyte	Total Number of Samples	Detection Frequency > Background ²⁵	Mean	Maximum Concentration	Standard Deviation	Background Mean Plus 2SD	Unit
439	Nickel	6	0.00%	7.67	13.1	3.27	14.8	mg/kg
440	Nickel	10	40.00%	12.8	21	5.46	14.8	mg/kg
441	Nickel	5	60.00%	16.2	20	3.11	14.8	mg/kg
442	Nickel	11	36.36%	12.8	19.1	4.20	14.8	mg/kg
444	Nickel	39	43.59%	13.8	21	3.92	14.8	mg/kg
445	Nickel	1	100.00%	21.0	21	NA	14.8	mg/kg
447	Nickel	17	76.47%	16.2	22	4.03	14.8	mg/kg
450	Nickel	1	100.00%	18.0	18	NA	14.8	mg/kg
455	Nickel	1	0.00%	5.10	5.1	NA	14.8	mg/kg
528	Nickel	1	100.00%	20.0	20	NA	14.8	mg/kg
559	Nickel	12	25.00%	11.7	25	5.74	14.8	mg/kg
701	Nickel	4	75.00%	15.0	19	3.74	14.8	mg/kg
707	Nickel	20	0.00%	6.43	13	2.26	14.8	mg/kg
712/713	Nickel	12	66.67%	17.7	45	11.7	14.8	mg/kg
770	Nickel	2	50.00%	16.0	20	5.66	14.8	mg/kg
771	Nickel	12	83.33%	18.0	31	6.49	14.8	mg/kg
774	Nickel	4	50.00%	13.7	16	3.35	14.8	mg/kg
776	Nickel	20	45.00%	14.8	37	7.48	14.8	mg/kg
777	Nickel	25	40.00%	18.5	77	18.3	14.8	mg/kg
778	Nickel	15	26.67%	12.1	22	4.99	14.8	mg/kg
779	Nickel	3	100.00%	20.0	22	2.65	14.8	mg/kg
865	Nickel	25	20.00%	10.7	23	5.00	14.8	mg/kg
881	Nickel	25	28.00%	15.3	56.7	9.20	14.8	mg/kg
883	Nickel	12	66.67%	15.9	22	5.07	14.8	mg/kg
886	Nickel	23	17.39%	11.4	18.4	3.47	14.8	mg/kg
889	Nickel	3	66.67%	18.3	24.6	6.15	14.8	mg/kg
991	Nickel	10	60.00%	14.4	27	7.86	14.8	mg/kg

NA = Not Applicable

2.1.16 Selenium

Selenium was not identified or discussed in building process information (CDH 1992; DOE 2004). Selenium has not been found associated with UBC sites (DOE 2004).

Selenium compounds were initially identified in the ChemRisk Task 1 report in inventory at RFETS (although no specific building was identified) in the form of dioxide, oxide, pellets, and powder. These chemicals appeared to have been used as laboratory standards or analytical testing materials because they were used in very small quantities (CDH 1991b). Based on the estimated quantity of these chemicals used, selenium was not carried forward as a material of concern for the ChemRisk process.

Small amounts of selenium waste have been generated from both laboratory and process buildings (Table 4 and Table 7).

Selenium was not identified above a RFCA AL requiring an accelerated action based on Closeout Reports for IHSSs and UBC sites.

In reviewing the selenium soil data beneath the slabs for UBC sites (Figure 16), the summary statistics presented in Table 23 were generated.

Table 23 UBC Soil Summary Statistics for Selenium (mg/kg)

Table 25 ODE Son Summary Statistics for Sciential (ing. kg)										
UBC	Analyte	Total Number of Samples	Detection Frequency > Background ²⁶	Mean	Maximum Concentration	Standard Deviation	Background Mean Plus 2SD	Unit		
122	Selenium	6	0.00%	ND	ND	ND	1.24	mg/kg		
123	Selenium	0	NA	NA	NA	NA	1.24	mg/kg		
331	Selenium	16	0.00%	ND	ND	ND	1.24	mg/kg		
371	Selenium	1	0.00%	ND	ND	ND	1.24	mg/kg		
374	Selenium	3	0.00%	ND	ND	ND	1.24	mg/kg		
439	Selenium	6	0.00%	ND	ND	ND	1.24	mg/kg		
440	Selenium	10	0.00%	ND	ND	ND	1.24	mg/kg		
441	Selenium	5	0.00%	ND	ND	ND	1.24	mg/kg		
442	Selenium	11	0.00%	ND	ND	ND	1.24	mg/kg		
444	Selenium	39	15.38%	0.741	2.1	0.442	1.24	mg/kg		
445	Selenium	1	100.00%	1.80	1.8	NA	1.24	mg/kg		
447	Selenium	17	0.00%	0.475	1.2	0.189	1.24	mg/kg		
450	Selenium	1	100.00%	1.30	1.3	NA	1.24	mg/kg		
455	Selenium	1	0.00%	ND	ND	ND	1.24	mg/kg		
528	Selenium	1	0.00%	ND	ND	ND	1.24	mg/kg		
559	Selenium	12	0.00%	ND	ND	ND	1.24	mg/kg		
701	Selenium	4	0.00%	ND	ND	ND	1.24	mg/kg		
707	Selenium	20	0.00%	ND	ND	ND	1.24	mg/kg		
712/713	Selenium	- 12	0.00%	0.388	0.45	0.101	1.24	mg/kg		
770	Selenium	2	0.00%	ND	ND	ND	1.24	mg/kg		
771	Selenium	12	0.00%	0.296	0.729	0.165	1.24	mg/kg		
774	Selenium	4	0.00%	ND	ND	ND	1.24	mg/kg		
776	Selenium	20	0.00%	ND	ND	ND	1.24	mg/kg		
777	Selenium	25	0.00%	ND	ND	ND	1.24	mg/kg		
778	Selenium	15	0.00%	0.432	0.88	0.125	1.24	mg/kg		
779	Selenium	3	0.00%	ND	ND	ND	1.24	mg/kg		
865	Selenium	25	0.00%	ND	ND	ND	1.24	mg/kg		
881	Selenium	25	0.00%	ND	ND	ND	1.24	mg/kg		
883	Selenium	12	0.00%	ND	ND	ND	1.24	mg/kg		
886	Selenium	23	0.00%	0.234	0.85	0.139	1.24	mg/kg		
889	Selenium	3	0.00%	ND	ND	ND	1.24	mg/kg		
991	Selenium	10	0.00%	0.495	1.1	0.215	1.24	mg/kg		

NA = Not Applicable

ND = Not Detected

2.1.17 Strontium

Strontium was not identified or discussed in building process information (CDH 1992; DOE 2004). Strontium has not been found associated with UBC sites (DOE 2004).

²⁶ This column indicates those samples that were detected and greater than background. As a result, if this field shows 0% and a mean and maximum concentration are reported, the values represent those analytes detected but less than background.

Strontium compounds were initially identified in the ChemRisk Task 1 report in inventory at RFETS (although no specific building was identified) in the form of carbonate, chloride, fluoride, nitrate, oxide, sulfide, and zirconate. These chemicals appeared to have been used as laboratory standards or analytical testing materials because they were used in very small quantities (CDH 1991b). Based on the estimated quantity of these chemicals used, strontium was not carried forward as a material of concern for the ChemRisk process.

There is no indication that strontium waste has been generated from on-site operations (Table 4 and Table 7).

Strontium was not identified above a RFCA AL requiring an accelerated action based on SAPs, SAP Addenda, or Closeout Reports for IHSSs and UBCs.

In reviewing the strontium soil data beneath the slabs for UBC sites (Figure 17), the summary statistics presented in Table 24 were generated.

Table 24 UBC Soil Summary Statistics for Strontium (mg/kg)

UBC	Analyte	Total Number of Samples	Detection Frequency > Background ²⁷	Mean	Maximum Concentration	Standard Deviation	Background Mean Plus 2SD	Unit
122	Strontium	6	0.00%	17.5	34	9.44	48.8	mg/kg
123	Strontium	0	NA	NA	NA	NA	48.8	mg/kg
331	Strontium	16	12.50%	28.9	78	17.5	48.8	mg/kg
371	Strontium	1	0.00%	8.30	8.3	NA	48.8	mg/kg
374	Strontium	3	0.00%	19.0	37.8	16.3	48.8	mg/kg
439	Strontium	6	0.00%	11.9	16.5	3.46	48.8	mg/kg
440	Strontium	10	0.00%	22.0	35	7.71	48.8	mg/kg
441	Strontium	5	0.00%	26.4	43	10.1	48.8	mg/kg
442	Strontium	11	27.27%	35.9	79	19.2	48.8	mg/kg
444	Strontium	39	10.26%	32.9	120	20.4	48.8	mg/kg
445	Strontium	1	0.00%	24.0	24	NA	48.8	mg/kg
447	Strontium	17	23.53%	36.5	81	18.1	48.8	mg/kg
450	Strontium	1	100.00%	60.0	60	NA	48.8	mg/kg
455	Strontium	1	0.00%	16.0	16	NA	48.8	mg/kg
528	Strontium	1	100.00%	97.0	97	NA	48.8	mg/kg
559	Strontium	12	8.33%	24.5	60	15.4	48.8	mg/kg
701	Strontium	4	25.00%	34.3	49	15.3	48.8	mg/kg
707	Strontium	20	0.00%	16.7	37	6.69	48.8	mg/kg
712/713	Strontium	12	33.33%	52.9	120	34.5	48.8	mg/kg
770	Strontium	2	100.00%	60.0	71	15.6	48.8	mg/kg
771	Strontium	12	41.67%	44.1	63.5	13.1	48.8	mg/kg
774	Strontium	4	75.00%	56.8	89	22.6	48.8	mg/kg
776	Strontium	20	0.00%	25.3	40	8.14	48.8	mg/kg
777	Strontium	25	16.00%	30.2	110	22.2	48.8	mg/kg
778	Strontium	15	0.00%	27.3	47	9.98	48.8	mg/kg
779	Strontium	3	0.00%	23.0	28	5.00	48.8	mg/kg

²⁷ This column indicates those samples that were detected and greater than background. As a result, if this field shows 0% and a mean and maximum concentration are reported, the values represent those analytes detected but less than background.

UBC	Analyte	Total Number of Samples	Detection Frequency > Background ²⁷	Mean	Maximum Concentration	Standard Deviation	Background Mean Plus 2SD	Unit
865	Strontium	25	48.00%	72.6	300	70.0	48.8	mg/kg
881	Strontium	25	24.00%	39.9	132	24.6	48.8	mg/kg
883	Strontium	12	91.67%	110	200	44.5	48.8	mg/kg
886	Strontium	23	47.83%	75.0	178	51.8	48.8	mg/kg
889	Strontium	3	100.00%	107	131	28.7	48.8	mg/kg
991	Strontium	10	40.00%	73.8	250	76.1	48.8	mg/kg

 $\overline{NA} = \overline{N}$ ot Applicable

2.1.18 Thallium

Thallium was not identified or discussed in building process information (CDH 1992; DOE 2004). Thallium has not been found associated in UBC sites (DOE 2004).

Thallium compounds were initially identified in the ChemRisk Task 1 report in inventory at RFETS (although no specific building was identified) (CDH 1991b). These chemicals appeared to have been used as laboratory standards or analytical testing materials because they were used in very small quantities. Based on the estimated quantity of these chemicals used, thallium was not carried forward as a material of concern for the ChemRisk process.

Small amounts of waste containing thallium as an underlying hazardous constituent have been generated from both laboratory and process buildings (Table 4 and Table 7).

Thallium was not identified above a RFCA AL requiring an accelerated action based on SAPs, SAP Addenda, or Closeout Reports for IHSSs and UBC sites.

In reviewing the thallium soil data beneath the slabs for UBC sites (Figure 18), the summary statistics presented in Table 25 were generated.

Table 25 UBC Soil Summary Statistics for Thallium (mg/kg)

				-			75 1 1	
UBC	Analyte	Total Number of Samples	Detection Frequency > Background ²⁸	Mean	Maximum Concentration	Standard Deviation	Background Mean Plus 2SD	Unit
122	Thallium	6	33.33%	0.681	1.3	0.373	0.443	mg/kg
123	Thallium	0	NA	NA	NA	NA	0.443	mg/kg
331	Thallium	16	12.50%	0.928	5.4	1.22	0.443	mg/kg
371	Thallium	1	0.00%	ND	ND	ND	0.443	mg/kg
374	Thallium	3	33.33%	0.445	0.91	0.403	0.443	mg/kg
439	Thallium	6	33.33%	0.568	0.93	0.272	0.443	mg/kg
440	Thallium	10	60.00%	0.821	1.2	0.324	0.443	mg/kg
441	Thallium	5	80.00%	0.980	1.1	0.130	0.443	mg/kg
442	Thallium	11	0.00%	ND	ND	ND	0.443	mg/kg
444	Thallium	39	41.03%	0.838	2	0.479	0.443	mg/kg
445	Thallium	1	0.00%	ND	ND	ND	0.443	mg/kg
447	Thallium	17	29.41%	0.790	2.1	0.569	0.443	mg/kg

²⁸ This column indicates those samples that were detected and greater than background. As a result, if this field shows 0% and a mean and maximum concentration are reported, the values represent those analytes detected but less than background.

UBC	Analyte	Total Number of Samples	Detection Frequency > Background ²⁸	Mean	Maximum Concentration	Standard Deviation	Background Mean Plus 2SD	Unit
450	Thallium	1	0.00%	ND	ND	ND	0.443	mg/kg
455	Thallium	1	0.00%	ND	ND	ND	0.443	mg/kg
528	Thallium	1	100.00%	1.80	1.8	NA	0.443	mg/kg
559	Thallium	12	16.67%	0.596	1.4	0.353	0.443	mg/kg
701	Thallium	4	0.00%	ND	ND	ND	0.443	mg/kg
707	Thallium	20	5.00%	0.485	1.2	0.169	0.443	mg/kg
712/713	Thallium	11	0.00%	ND	ND	ND	0.443	mg/kg
770	Thallium	2	0.00%	ND	ND	ND	0.443	mg/kg
771	Thallium	13	23.08%	0.379	1.2	0.287	0.443	mg/kg
774	Thallium	4	0.00%	0.490	0.37	0.085	0.443	mg/kg
776	Thallium	20	15.00%	0.583	1.4	0.300	0.443	mg/kg
777	Thallium	25	8.00%	0.481	1.1	0.175	0.443	mg/kg
778	Thallium	15	6.67%	0.510	1.3	0.219	0.443	mg/kg
779	Thallium	3	0.00%	ND	ND	ND	0.443	mg/kg
865	Thallium	25	28.00%	0.621	1.1	0.269	0.443	mg/kg
881	Thallium	25	0.00%	ND	ND	ND	0.443	mg/kg
883	Thallium	12	66.67%	1.62	5.1	1.25	0.443	mg/kg
886	Thallium	23	8.70%	0.343	1	0.208	0.443	mg/kg
889	Thallium	3	0.00%	ND	ND	ND	0.443	mg/kg
991	Thallium	10	0.00%	ND	ND	ND	0.443	mg/kg

NA = Not Applicable

ND = Not Detected

2.1.19 Vanadium

Pit construction in Building 707 generally used plutonium, uranium, beryllium, aluminum, and stainless steel (CDH 1992; DOE 2004). However, in some instances more exotic materials such as cadmium, vanadium, silver, and gold were used; however, the amounts were relatively minor compared to the primary five metals. The metallurgical operations in Building 865 (R&D) involved the development of alloys in the 1970s. Some of the metals employed in the alloying development included aluminum, copper, magnesium, molybdenum, niobium, platinum, stainless steel, tantalum, titanium, and vanadium. Vanadium was also identified as associated with metalworking in Building 444. In Building 447 materials handled included stainless steel, beryllium, aluminum, depleted uranium, and vanadium compounds.

The buildings identified above involved radiological operations and included extensive HEPA filtration systems. Any emissions from machining vanadium would have been collected on these filters prior to release from the buildings.

Vanadium compounds were initially identified in the ChemRisk Task 1 report in inventory at RFETS (CDH 1991b). However, based on the estimated quantity of these chemicals used (typically less than 1 kg with the exception of a pentoxide at 12 kg in 1974 and less than 1 kg in 1988) vanadium was not carried forward as a material of concern for the ChemRisk process.

Waste containing vanadium as an underlying hazardous constituent was generated from Buildings 559 and 881 (Table 4 and Table 7).

Vanadium was not identified above a RFCA AL requiring an accelerated action based on SAPs, SAP Addenda, or Closeout Reports for IHSSs and UBC sites.

In reviewing the vanadium soil data beneath the slabs for UBC sites (Figure 19), the summary statistics presented in Table 26 were generated.

Table 26 UBC Soil Summary Statistics for Vanadium (mg/kg)

		Total	Detection		<u> </u>		Background	
UBC	Analyte	Number of	Frequency	Mean	Maximum	Standard	Mean Plus	Unit
		Samples	> Background ²⁹		Concentration	Deviation	2SD	
122	Vanadium	6	0.00%	24.2	42	13.3	43.1	mg/kg
123	Vanadium	0	NA	NA	NA	NA	43.1	mg/kg
331	Vanadium	16	6.25%	30.8	50	7.43	43.1	mg/kg
371	Vanadium	1	0.00%	12.0	12	NA	43.1	mg/kg
374	Vanadium	3	0.00%	16.9	28.7	10.5	43.1	mg/kg
439	Vanadium	6	0.00%	17.8	35.5	9.20	43.1	mg/kg
440	Vanadium	10	0.00%	26.1	39	8.76	43.1	mg/kg
441	Vanadium	5	20.00%	39.2	52	8.07	43.1	mg/kg
442	Vanadium	11	18.18%	34.8	46.3	8.75	43.1	mg/kg
444	Vanadium	39	7.69%	33.3	51	8.36	43.1	mg/kg
445	Vanadium	1	100.00%	54.0	54	NA	43.1	mg/kg
447	Vanadium	17	17.65%	39.7	52	8.59	43.1	mg/kg
450	Vanadium	1	0.00%	31.0	31	NA	43.1	mg/kg
455	Vanadium	1	0.00%	21.0	21	NA	43.1	mg/kg
528	Vanadium	1	100.00%	78.0	78	NA	43.1	mg/kg
559	Vanadium	12	0.00%	24.0	42	8.98	43.1	mg/kg
701	Vanadium	4	0.00%	35.5	43	8.35	43.1	mg/kg
707	Vanadium	20	0.00%	15.1	26	3.98	43.1	mg/kg
712/713	Vanadium	12	16.67%	32.6	74	17.4	43.1	mg/kg
770	Vanadium	2	0.00%	35.5	39	4.95	43.1	mg/kg
771	Vanadium	12	8.33%	30.2	48	8.76	43.1	mg/kg
774	Vanadium	4	0.00%	34.8	41	4.19	43.1	mg/kg
776	Vanadium	20	15.00%	29.0	51	12.2	43.1	mg/kg
777	Vanadium	25	8.00%	25.8	56	11.3	43.1	mg/kg
778	Vanadium	15	13.33%	26.2	44	9.96	43.1	mg/kg
779	Vanadium	3	33.33%	39.0	53	14.5	43.1	mg/kg
865	Vanadium	25	8.00%	27.8	46	9.02	43.1	mg/kg
881	Vanadium	25	4.00%	32.8	49.3	6.66	43.1	mg/kg
883	Vanadium	12	66.67%	47.3	64	15.0	43.1	mg/kg
886	Vanadium	23	4.35%	24.7	64	10.9	43.1	mg/kg
889	Vanadium	3	66.67%	59.1	76.9	22.7	43.1	mg/kg
991	Vanadium	10	50.00%	39.7	70	21.1	43.1	mg/kg

NA = Not Applicable

²⁹ This column indicates those samples that were detected and greater than background. As a result, if this field shows 0% and a mean and maximum concentration are reported, the values represent those analytes detected but less than background.

2.1.20 Zinc

Zinc was not identified or discussed in building process information (CDH 1992; DOE 2004). Zinc has not been found associated with UBC sites (DOE 2004).

Zinc was initially identified in the ChemRisk Task 1 report in inventory at RFETS (although no specific building was identified) (CDH 1991b), in the form of acetate, bromide, carbonate, chloride, cyanide, fluoride, metal powder, nitrate, oxide, sulfide, and sulfate. Based on the relative toxicity of the material, how the material might have been released into the environment, and/or the likelihood for transport off-site, zinc was not carried forward as a material of concern for the ChemRisk process.

Waste containing zinc as an underlying hazardous constituent was generated from Building 881 (Table 4 and Table 7).

Zinc was not identified above a RFCA AL requiring an accelerated action based on SAPs, SAP Addenda, or Closeout Reports for IHSSs and UBC sites.

Zinc orthophosphate was added to the drinking water system from 2002 to system closure to prevent copper and lead corrosion.

In reviewing the zinc soil data beneath the slab for UBC sites (Figure 20), the summary statistics presented in Table 27 were generated.

Table 27 UBC Soil Summary Statistics for Zinc (mg/kg)

UBC	Analyte	Total Number of Samples	Detection Frequency > Background ³⁰	Mean	Maximum Concentration	Standard Deviation	Background Mean Plus 2SD	Unit
122	Zinc	6	0.00%	28.3	45	11.7	74.2	mg/kg
123	Zinc	0	NA	NA	NA	NA	74.2	mg/kg
331	Zinc	16	6.25%	35.6	190	42.3	74.2	mg/kg
371	Zinc	1	0.00%	14.0	14	NA	74.2	mg/kg
374	Zinc	3	0.00%	27.2	47.6	17.9	74.2	mg/kg
439	Zinc	6	16.67%	59.0	183	60.8	74.2	mg/kg
440	Zinc	10	0.00%	33.7	45	9.31	74.2	mg/kg
441	Zinc	5	20.00%	47.6	120	40.6	74.2	mg/kg
442	Zinc	11	0.00%	32.4	69.6	15.1	74.2	mg/kg
444	Zinc	39	2.56%	34.6	130	19.1	74.2	mg/kg
445	Zinc	1	0.00%	28.0	28	NA	74.2	mg/kg
447	Zinc	17	11.76%	48.5	270	62.4	74.2	mg/kg
450	Zinc	1	0.00%	25.0	25	NA	74.2	mg/kg
455	Zinc	1	100.00%	80.0	80	NA	74.2	mg/kg
528	Zinc	1	0.00%	42.0	42	NA	74.2	mg/kg
559	Zinc	12	0.00%	21.2	37	6.83	74.2	mg/kg
701	Zinc	4	0.00%	51.5	69	15.4	74.2	mg/kg
707	Zinc	20	0.00%	28.0	45	8.38	74.2	mg/kg

³⁰ This column indicates those samples that were detected and greater than background. As a result, if this field shows 0% and a mean and maximum concentration are reported, the values represent those analytes detected but less than background.

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UBC	Analyte	Total Number of Samples	Detection Frequency > Background ³⁰	Mean	Maximum Concentration	Standard Deviation	Background Mean Plus 2SD	Unit
712/713	Zinc	12	33.33%	112	410	116	74.2	mg/kg
770	Zinc	2	0.00%	49.0	64	21.2	74.2	mg/kg
771	Zinc	12	16.67%	53.2	93.6	20.7	74.2	mg/kg
774	Zinc	4	0.00%	48.0	59	7.87	74.2	mg/kg
776	Zinc	20	0.00%	35.2	60	9.59	74.2	mg/kg
777	Zinc	25	4.00%	39.5	160	28.8	74.2	mg/kg
778	Zinc	15	0.00%	33.1	56	10.0	74.2	mg/kg
779	Zinc	3	0.00%	26.7	29	2.52	74.2	mg/kg
865	Zinc	25	8.00%	49.9	180	31.0	74.2	mg/kg
881	Zinc	25	0.00%	35.1	72.7	16.2	74.2	mg/kg
883	Zinc	12	0.00%	45.8	63	12.2	74.2	mg/kg
886	Zinc	23	4.35%	31.4	124	22.4	74.2	mg/kg
889	Zinc	3	0.00%	48.2	66.6	18.5	74.2	mg/kg
991	Zinc	10	0.00%	41.1	74	16.4	74.2	mg/kg

NA = Not Applicable

2.2 Radionuclides

A total of 4 radionuclide AOIs have been identified in media based on the nature and extent evaluations. These radionuclides are discussed in the following sections.

2.2.1 Cesium-137

The ChemRisk Task 1 Report identified cesium-137 as a radionuclide used for research, analytical, and calibration activities (for example, sealed and plated sources) (CDH 1991a). Based on the limited quantity of this material, cesium-137 was not carried forward through the ChemRisk process (CDH 1991b). In addition, the only cesium-137 waste generated at this site was identified as sealed sources (based on WEMS and WSRIC) from Buildings707, 776/777, and 991.

In addition, according to the Task 2 ChemRisk Report, environmental sampling data indicate the presence of detectable quantities of other radionuclides characteristic of nuclear weapons fallout, such as strontium-89 and 90, zirconium-95, cesium-137, and cerium-144, which were also found in environmental samples from 1970 through 1981. Detection of these compounds is consistent with the presence of fission products from worldwide fallout, and the detected levels are typical of other sites sampled in the western United States (CDH 1991b).

Based on a study of off-site areas surrounding RFETS, the Citizen's Environmental Sampling Committee (CESC) conducted soil and sediment sampling in 1993 and 1994, and analyzed for plutonium (plutonium-238 and plutonium-239/240), americium (americium-241), cesium (cesium-137), strontium (strontium -90), and uranium (uranium-235 and uranium-238) (CESC 1996). Background levels of cesium-137 and strontium-90 were detected in some soil samples. This report noted that cesium-137 and strontium-90 are generally associated with nuclear chain reactions. Although the Rocky Flats Plant never operated a full-scale nuclear reactor, they did perform criticality experiments. This report concluded that "no evidence has been found to suggest that cesium-137 or strontium-90 were released during the operational period of the Rocky Flats Plant".

Based on the history of usage and historical conclusions made regarding cesium-137, only two surface soil samples were collected during accelerated actions (one east of building 454 and one south of building 447), and both were below background (1.69 pCi/g). As a result, no summary statistics or figures were generated.

2.2.2 Radium

The ChemRisk Task 1 Report identified radium-226 as a radionuclide used for research, analytical, and calibration activities (for example, sealed and plated sources) (CDH 1991a). Based on the limited quantity of this material, radium-226 was not carried forward through the ChemRisk process (CDH 1991b). In addition, the only radium-226 waste generated at RFETS was identified as sealed sources (based on WEMS and WSRIC).

Radium-226 waste (sealed sources) was generated from Buildings 707, and 776/777.

The ChemRisk Task 1 Report did not identify radium-228 as a radionuclide used at the Rocky Flats Plant (CDH 1991a) and no radium-228 waste was reported to have been generated.

Radium-226 soil data around buildings and UBC sites are presented on Figure 21. Because there were no UBC soil data for radium-226, no summary statistics were generated.

Radium-228 soil data around buildings and UBC sites are presented on Figure 22. Because there were no UBC soil data for radium-228, no summary statistics were generated. For additional information regarding radium-228 see Section 2.2.4.

2.2.3 Strontium-89/90

The ChemRisk Task 1 Report identified stronium-89/90 as a radionuclide used for research, analytical, and calibration activities (for example, sealed sources, plated sources, liquid sources, and analytical stock solutions (CDH 1991a). Based on the limited quantity of this material, strontium-89/90 was not carried forward through the ChemRisk process (CDH 1991b).

In addition, according to the ChemRisk Task 2 Report, environmental sampling data indicate the presence of detectable quantities of other radionuclides characteristic of nuclear weapons fallout, such as strontium-89 and 90, zirconium-95, cesium-137, and cerium-144, which were also found in environmental samples from 1970 through 1981. Detection of these compounds is consistent with the presence of fission products from worldwide fallout, and the detected levels are typical of other sites sampled in the western United States (CDH 1991b).

Based on a study of off-site areas surrounding Rocky Flats, the CESC conducted soil and sediment sampling in 1993 and 1994 and analyzed for plutonium (Pu-238 and Pu-239/240), americium (americium-241), cesium (cesium-137), strontium (strontium-90), and uranium (uranium-235 and uranium-238) (CESC 1996). Background levels of cesium-137 and strontium-90 were detected in some soil samples. This report noted that cesium-137 and strontium-90 are generally associated with nuclear chain reactions. Although the Rocky Flats Plant never operated a full-scale nuclear reactor, they did perform criticality experiments. This

report concluded that "no evidence has been found to suggest that cesium-137 or strontium-90 were released during the operational period of the Rocky Flats Plant".

Based on the history of usage and historical conclusions made regarding strontium-89/90, only three soil samples were collected during accelerated actions for UBC 123. These results indicated that strontium 89/90 was not detected. As a result, no summary statistics or figure was generated.

2.2.4 Thorium-232

Thorium-232 has been identified in this review because the AOI radium-228 is in the thorium-232 decay chain.

Thorium has been used in several ways at RFETS since 1952. The major use was fabrication of metal parts from natural thorium and thorium alloys (Building 881). The compounds have been used in analytical procedures and development programs (Building 771). Although amounts were small, applications were numerous. Over the period from 1952 to 1976, which saw the majority of thorium applications at Rocky Flats, the quantity of thorium that was present varied from none to approximately 238 kg in any one month (CDPHE 1994).

A project in Building 881 involved thorium-232 production over several years in the late 1950s to early 1960s (CDH 1992). There were very tight controls, and thorium went through the same processes as enriched uranium; however, most was sent to Savannah River, South Carolina or Oak Ridge, Tennessee for recovery.

According to the March 1994 Task 5 Report, a large fraction of the thorium material discarded (that is, Normal Operational Loss) would likely have been in solid wastes or particulates trapped by ventilation exhaust filters, given that the primary use of Rocky Flats' thorium was for metalworking processes (CDPHE 1994). This report concludes that it is likely that less than 32 kg of thorium would have been released in airborne effluents from the Rocky Flats Plant over its operational history. In addition, criticality experiments were not likely to have been a source of significant releases of radionuclides from the Plant.

The information reviewed in the ChemRisk reports concludes that thorium-232 has not been a significant component of airborne effluents from the Rocky Flats Plant and was not used in significant quantities relative to other production radionuclides (CDH 1991b). Because thorium would most likely have been emitted in particulate form, and thorium-232 emits an alpha particle with each decay, thorium emissions are reflected in results of measurements of total long-lived alpha radioactivity that were performed since the early 1950s. Thorium operations have been insignificant relative to the primary production activities centered around plutonium and uranium, and little data exist to support the quantification of release. Therefore, efforts to attribute a portion of total long-lived alpha activity measurements to thorium-232 were not made in the ChemRisk reports (CDPHE 1994). In addition, because of the apparent diminishing of applications of thorium compounds since the 1970s, development of source term estimates for thorium-232 during the 1970s and 1980s was not considered warranted in the ChemRisk reports.

Based on the history of usage and historical conclusions made regarding thorium-232, no soil samples were collected during accelerated actions. As a result, no summary statistics or figure for thorium-232 was generated.

3.0 SUMMARY

Historical knowledge indicates there is no inventory remaining at RFETS for these AOIs, and any potential "source term" remaining is expected to be low, because the release potential was also low.

Table 28 provides a summary of information available for each AOI as discussed in previous sections.

4.0 REFERENCES

CDH, 1991a, Project Task 1 Report (Revised 1) Identification of Chemicals and Radionuclides Used at Rocky Flats, Colorado Department of Health prepared by ChemRisk, March.

CDH, 1991b, Project Task 2 Selection of the Chemicals and Radionuclides of Concern, Colorado Department of Health prepared by ChemRisk, June.

CDH, 1992, Project Tasks 3 & 4 Final Draft Report, Reconstruction of Historical Rocky Flats Operations & Identification of Release Points, Colorado Department of Health prepared by ChemRisk, August.

CDPHE 1994 Project Task 5 Report, Estimating Historical Emissions from Rocky Flats 1952-1989, Colorado Department of Public Health and Environment prepared by ChemRisk, March.

CESC, 1996, Soil and Sediment Study of Off-Site Areas Surrounding the Rocky Flats Nuclear Weapons Plant, Executive Summary, Citizen's Environmental Sampling Committee, September. Found at http://www.cdphe.state.co.us/rf/soilsedstudy.htm.

DOE, 1992, Final Historical Release Report as modified for Rocky Flats Plant, Golden, Colorado, June.

DOE, 2003a, Final Industrial Area Sampling and Analysis Plan FY03 Addendum Number IA-03-01 IHSS Groups 300-3, 300-4, 400-8, 700-4, 800-1, and 900-3, January. Approved by CDPHE on January 14, 2003.

DOE, 2003b, Final Proposed Action Memorandum for IHSS 101 and RCRA Closure of the RFETS Solar Evaporation Ponds, April. Approved by CDPHE on May 22, 2003.

DOE, 2004a, Industrial Area and Buffer Zone Sampling and Analysis Plan, Rocky Flats Environmental Technology Site, Golden, Colorado, May.

DOE, 2004b, Final Comprehensive Risk Assessment Work Plan and Methodology, September.

Table 28 Summary of AOI Information

AOIs	Examples of Materials	Inventory	Total	Waste	Spills/ Releases	IHSS, PAC or	UBC Mean Data	UBC Mean Data
71015	Used	in	Quantity in	Generated	Within	UBC Requiring	Results >	Results <
		Buildings	Inventory	in Buildings	Buildings ^b	an Accelerated	Background Mean +	Background Mean +
		Dunamgo	(kg) ^a	in Dundings	2	Action	2 Standard	2 Standard
			()				Deviations	Deviations
Aluminum	Aluminum metal,	444, 447,	7,700	371, 559,	None identified	None	441, 442, 445, 447,	122, 123, 331, 371,
	nitrate, silicate, and	707, 771,	,	707, 778,			450, 528, 712/713,	374, 439, 440, 444,
	oxide.	779, 883,		779, and 887			774, and 779	455, 559, 701, 707,
		and 865						770, 771, 776, 777,
							Ļ	778, 865, 881, 883,
								886, 889, and 991
Antimony	Acid containing	None	8	559°	None identified	None	712/713	122, 123, 331, 371,
	antimony, antimony	identified						374, 439, 440, 441,
	iodide, oxide,							442, 444, 445, 447,
	pentachloride, powder,					}		450, 455, 528, 559,
	trioxide, and trichloride.							701, 707, 770,
								771,774, 776, 777,
								778, 779, 865, 881,
								883, 886, 889, and 991
Arsenic	Arsenic acid, iodide,	None	5	371, 374,	None identified	PAC 700-137,	771	122, 123, 331, 371,
	metals, pentoxide,	identified		460, 559,		outside Building		374, 439, 440, 441,
	solution 3103, trioxide,			707, 771,		707, and PAC		442, 444, 445, 447,
	and oxide.			774,		SE-1602		450, 455, 528, 559,
	ļ			776/777,				701, 707, 712/713,
				778, 779,				770, 774, 776, 777,
				881, 883,				778, 779, 865, 881,
				865, 887,				883, 886, 889, and 991
			22	and 889	37 11 10 1	,,	500 1554	100 100 221 251
Barium	Barium acetate,	None	33	371, 374,	None identified	None	528 and 774	122, 123, 331, 371,
	chloride, fluoride,	identified		439, 442,				374, 439, 440, 441,
	metal, nitrate, and			444, 559,				442, 444, 445, 447,
	sulfate.			707, 771,				450, 455, 559, 701,
				774,				707, 712/713, 770,
				776/777,				771, 776, 777, 778,
			:	778, 779,				779, 865, 881, 883,
				881, 883,				886, 889, and 991
				865, 886, and 887				
Cadmium	Cadmium acetate,	122 444	44	122, 123,	123, 444, 528,	None	None	122, 123, 331, 371,
Cadmium	1	123, 444,	44	122, 123, 125, 331,	123, 444, 328, 559, 707, 731,	none	None	1
	chloride, iodide, metal,	528, 559,			771, and 774			374, 439, 440, 441,
	nitrate, and sulfate	561, 707,		371, 374, 439, 440,	//1, and //4			442, 444, 445, 447,
		731, 774,		439, 440,			1	450, 455, 528, 559,

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AOIs	Examples of Materials Used	Inventory in Buildings	Total Quantity in Inventory (kg) ^a	Waste Generated in Buildings	Spills/ Releases Within Buildings ^b	IHSS, PAC or UBC Requiring an Accelerated Action	UBC Mean Data Results > Background Mean + 2 Standard Deviations	UBC Mean Data Results ≤ Background Mean + 2 Standard Deviations
		771, 776/777, 881, and 883		441, 442, 444, 447, 460, 559, 701, 707, 771, 774, 776, 777, 778, 779, 865, 881, 883, 886, 887, and 991				701, 707, 712/713, 770, 771, 774, 776, 777, 778, 779, 865, 881, 883, 886, 889, and 991
Chromium (total)	Chromium boride, chloride, metal, nitrate, oxide, sulfate, and trioxide.	123, 371, 374, 444, 460, 528, 559, 561, 712/713, 731, 774, 776/777, and 779	734	122, 123, 331, 371, 374, 439, 440, 441, 444, 447, 460, 559, 701, 707, 771, 774, 776, 777, 778, 779, 865, 881, 883, 886, 887, and 991	123, 371, 374, 444, 528, 559, 707, 731, and 774	PAC 500-158	331, 440, 441, 444, 445, 447, 528, 701, 712/713, 770, 771, 774, 776, 777, 779, and 881	122, 123, 371, 374, 439, 442, 450, 455, 559, 707, 712/713, 778, 865, 883, 886, 889, and 991
Cobalt	Cobalt (metal, powder, wire, and foil), chloride, nitrate, oxide, and sulfate.	None identified	25	None identified	None identified	None	444, 450, 776, 777, 778, 779, 881, and 991	122, 123, 331, 371, 374, 439, 440, 441, 442, 445, 447, 455, 528, 559, 701, 707, 712/713, 770, 771, 774, 865, 883, 886, and 889
Copper	Copper metal, cyanide and sulfate. Ferric ammonium	444, 779, 881, 883, and 865	978	559 and 881	None identified None identified	None	331, 374, 440, 444, 447, 450, 455, 528, 559, 701, 707, 712/713, 770, 771, 774, 776, 777, 778, 779, 865, 881, 883, 886, 889, and 991 445, 447, 712/713,	122, 123, 371, 439, 441, 442, and 445

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AOIs	Examples of Materials Used oxalate, nitrate, sulfide, and ammonium sulfate.	Inventory in Buildings 445, 450, and 455	Total Quantity in Inventory (kg) ²	Waste Generated in Buildings identified	Spills/ Releases Within Buildings ^b	IHSS, PAC or UBC Requiring an Accelerated Action	UBC Mean Data Results > Background Mean + 2 Standard Deviations 779, 883, and 889	UBC Mean Data Results ≤ Background Mean + 2 Standard Deviations 374, 439, 440, 441, 442, 444, 450, 455, 528, 559, 701, 707,
								770, 771, 774, 776, 777, 778, 865, 881, 886, and 991
Lead	Lead acetate, chloride, iodide, metal, nitrate, oxide and powder.	123, 371, 374, 528, 559, 701, 707, 731, 774, 771, and 776/777	140	122, 123, 125, 331, 371, 374, 439, 440, 442, 447, 460, 559, 701, 707, 770, 771, 774, 776/777, 778, 779, 865, 881, 883, 886, 887, 889, and 991	123, 371, 374, 528, 559, 707, 731,771, and 774	UBC 123, PAC 400-122, PAC NW-1505, and PAC SE-1602	447	122, 123, 331, 371, 374, 439, 440, 441, 442, 444, 445, 450, 455, 528, 559, 701, 707, 712/713, 770, 771, 774, 776, 777, 778, 779, 865, 881, 883, 886, 889, and 991
Lithium	Lithium aluminum hydride, chloride, chromate, fluoride, metal, nitrate, and sulfate.	559, 707, 776/777, and 881	300	125, 371, 444, 460, 559, 771, 778, 881, and 883	559 and 707	None	441, 445, 447, 712/713, 774, 776, 779, 881, 883,	122, 123, 331, 371, 374, 439, 440, 442, 444, 450, 455, 528, 559, 701, 707, 770, 771, 777, 778, 865, 886, 889, and 991
Manganese	Manganese carbonate, chips, dioxide, flake, II oxide, metal, powder, chloride, nitrate and sulfate.	None identified	6	None identified	None identified	None	883 and 889	122, 123, 331, 371, 374, 439, 440, 441, 442, 444, 445, 447, 450, 455, 528, 559, 701, 707, 712/713, 770, 771, 774, 776, 777, 778, 779, 865, 881, 886, and 991
Mercury	Mercuric acetate, chloride, mercuric oxide.		10					

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AOIs	Examples of Materials Used	Inventory in Buildings	Total Quantity in Inventory (kg) ^a	Waste Generated in Buildings	Spills/ Releases Within Buildings ^b	IHSS, PAC or UBC Requiring an Accelerated Action	UBC Mean Data Results > Background Mean + 2 Standard Deviations	UBC Mean Data Results ≤ Background Mean + 2 Standard Deviations
Molybdenum	Molybdenum boride, carbide, disulfide, metal, metal powder, silicide, and trioxide.	444, 771, 774, 776/777, 779, and 881	13	559	None identified	None	712/713 and 883	122, 123, 331, 371, 374, 439, 440, 441, 442, 444, 445, 447, 450, 455, 528, 559, 701, 707, 770, 771, 774, 776, 777, 778, 779, 865, 881, 886, 889, and 991
Nickel	Nickel acetate, chloride, cyanide, metal, metal powder, nitrate, oxide and sulfate	371, 374, 444, 771, 774, 776/777, 779, 881, and 991	126	122, 125, 331, 371, 374, 439, 442, 444, 559, 701, 707, 771, 774, 776/777, 778, 779, 865, 881, 883,886, and	371, 374, 444, 771, and 774	None	331, 441, 445, 447, 450, 528, 701, 712/713, 770, 771, 777, 779, 881, 883, and 889	122, 123, 371, 374, 439, 440, 442, 444, 447, 455, 559, 707, 774, 776, 778, 865, 886, and 991
Selenium	Selenium dioxide, oxide, pellets, and powder	None identified	1	371, 374, 439, 441, 444, 559, 707, 771, 774, 776/777, 779, 865, 881, 883, and 887	None identified	None	445 and 450	122, 123, 331, 371, 374, 439, 440, 441, 442, 444, 447, 455, 528, 559, 701, 707, 712/713, 770, 771, 774, 776, 777, 778, 779, 865, 881, 883, 886, 889, and 991
Strontium	Strontium carbonate, chloride, fluoride, nitrate, oxide, sulfide and zirconate	None identified	8	None identified	None identified	None	450, 528, 712/713, 770, 774, 865, 883, 886, 889, and 991	122, 123, 331, 371, 374, 439, 440, 441, 442, 444, 445, 447, 455, 559, 701, 707, 771, 776, 777, 778, 779, and 881
Thallium	Thallium	None identified	<1	559, 707, 771, 779, and 881	None identified	None	122, 331, 374, 439, 440, 441, 444, 447, 528, 559, 707, 774,	123, 371, 442, 445, 450, 455, 701, 712/713, 770, 771,

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AOIs	Examples of Materials Used	Inventory in Buildings	Total Quantity in Inventory (kg) ^a	Waste Generated in Buildings	Spills/ Releases Within Buildings ^b	IHSS, PAC or UBC Requiring an Accelerated Action	UBC Mean Data Results > Background Mean + 2 Standard Deviations 776, 777, 778, 865,	UBC Mean Data Results ≤ Background Mean + 2 Standard Deviations 779, 881, 889, and 991
							883, and 886	775, 001, 005, und 551
Vanadium	Vanadium carbide, metal, nitride, pentoxide and sulfate	444, 447, 707, 779, 865, 881, 883,	4	559 and 881	None identified	None	445, 528, 883, and 889	122, 123, 331, 371, 374, 439, 440, 441, 442, 444, 447, 450, 455, 559, 701, 707, 712/713, 770, 771, 774, 776, 777, 778, 779, 865, 881, 886, and 991
Zinc	Zinc acetate, bromide, carbonate, chloride, chromate primer, cyanide, fluoride, metal powder, mossy, nitrate, oxide and sulfate	None identified	86	881	None identified	None	455	122, 123, 331, 371, 374, 439, 440, 441, 442, 444, 445, 447, 450, 528, 559, 701, 707, 712/713, 770, 771, 774, 776, 777, 778, 779, 865, 881, 883, 886, 889, and 991
Cesium-137	Sealed and plated sources	886	No quantity specified	707, 776/777, and 991	None identified	None	All samples collected were below background	NA
Radium-226	Sealed and plated sources	None identified	No quantity specified	707 and 776/777	None identified	None	NA	NA
Radium-228 (Thorium- 232)	Radium-228 was not identified in inventory at RFETS. Thorium-232 was used	771 and 881	No quantity specified	881	None identified	None	NA	NA
Strontium- 89/90	Sealed, plated and liquid sources, and analytical stock solutions	779	No quantity specified	707 and 991	None identified	None	All samples collected were not detected	NA

NA = Not Applicable because data was not collected.

^a Based on a 1988 inventory in the ChemRisk Task One Report (CDH 1991a)

^b No spills or releases of these AOIs occurred within a building that required an action prior to demolition, based on sampling and analysis results.

^c Antimony was identified as an underlying hazardous constituent in a waste from this building.

DOE, 2004c, Closeout Report for IHSS Group 500-2, IHSS 500-158 Radioactive Site-Building 551, June. Approved by CDPHE on June 18, 2004.

DOE, 2005a, Background Data Summary Tables Attachment 2, Data Description and Evaluation Volume 2 of 15, Appendix A Comprehensive Risk Assessment of the Draft Remedial Investigation/Feasibility Study Report, September.

DOE, 2005b, Closeout Report for IHSS Group 900-11, PAC SE-1602 East Firing Range and Target Area, March. Approved by EPA, Region VIII on February 8, 2005.

KH, 1998, Building 771/774 Cluster Closure Project RLCR, Rev. 2, (Administrative Record # B771-A-000119), Kaiser-Hill Company, L.L.C., Rocky Flats Environmental Technology Site, Golden, Colorado, August 8.

RFO, 1993, ORPS Occurrence Report Number: RFO--KHLL-371OPS-2003-0019, Rocky Flats Environmental Technology Site, Golden, Colorado, June 30.

RFO, 1998, ORPS Occurrence Report Number: RFO--KHLL-LIQWASTE-1998-0002, Rocky Flats Environmental Technology Site, Golden, Colorado, June 9.

RFO, 2002, ORPS Occurrence Report Number: RFO--KHLL-374OPS-2002-0004, Rocky Flats Environmental Technology Site, Golden, Colorado, November 25.

These occurrence reports were taken from the RFETS database Occurrence Reporting and Processing System (ORPS), which identified emergencies from 1991 to site closure.

Occurrence Report Number: RFO--KHLL-371OPS-2003-0019

On June 30, 1993 there was a sprinkler head malfunction in Room 3189 of Building 371 (radiological material area [not a contamination area or a radiological buffer area]), resulting in the release of fire water that ran into the hallway and into rooms 3187 A & B and 33185, eventually running onto Dock 18T and onto the ground north of Building 374 towards the outside storm drain. Fire department personnel began diverting water away from the storm drain. Approximately 6,000 gallons of water was released. It was collected and sampled and determined to be clean and approved for release to the storm drain system. (per emergency reporting system)

Impact on Environment, Safety, and Health: Radiological surveys and samples from monitoring equipment verified that this event caused no release of radiological materials. The environment and the health and safety of the public and plant personnel were not threatened.

Occurrence Report Number: RFO--KHLL-LIQWASTE-1998-0002

On June 9, 1998, at 1030 hours, approximately 1 gallon of a dark green liquid was discovered on Dock 8 of Buildings 371/374. Building 374 Environmental Operations and Radiological Operations personnel were contacted to investigate the substance. The substance was determined to be phosphoric acid contaminated with depleted uranium. The phosphoric acid apparently came from the drain pipe for Tank D-843. Radiological surveys taken indicated levels up to 3,000 direct counts per minute of alpha contamination. Incident Command was immediately established by the Building 371 Shift Manager. The RFETS Shift Superintendent responded and assumed command over the incident and declared an Operational Emergency at the Alert-Star level. This resulted in the precautionary activation of the Emergency Operations Center.

Tank D-843 had been filled with the phosphoric acid solution to the point that the high-level alarm had been actuated since 1991. The trigger for this event was the emergency generator test conducted on June 8, 1998, which de-energized the vent scrubber system, a vacuum system designed to remove fumes above the acid in the tank. The tank had become completely filled with liquid over the years and, consequently, the vacuum of the vent scrubber had drawn liquid up into the vent line. When the vent scrubber was de-energized, this vacuum was lost. Liquid in the dock drain line (the only input line to the tank), which had also become full, was then forced out onto the dock.

Impact on Environment, Safety, and Health: Although radioactive/toxic material was spilled in an undesirable location, it was determined that there was no impact on the environment, or on the health or safety of workers or the public.

RFO 2002

Occurrence Report Number: RFO--KHLL-374OPS-2002-0004

On November 25, 2002, there was a spill of low-level mixed waste from the RCRA-regulated Tank T231A (located south of Buildings 371/374) sludge removal operation. The spill occurred during a compressed air blow down of the 4-inch hose line between the pump and the centrifuge following completion of the sludge removal operations. The purpose of the system blow down was to empty the line of sludge prior to freezing weather conditions. During the blow down of the system the 4-inch line separated from its connection near the centrifuge causing the spill condition.

The spill exceeded the reportable quantity (10 pounds) for an F-listed waste. The majority of the waste was released inside the secondary containment area for the tank. Approximately 1 to 5 gallons (8 to 42 pounds) of waste was released outside the secondary containment over an area of approximately 600 square feet. Approximately 10 to 25 gallons (83 to 210 pounds) of waste was released inside of secondary containment over an area of approximately 200 square feet.

The spill was contained and the highest reported contamination levels were up to 750 disintegrations per minute per 100 square centimeters (dpm/100 cm²) both inside and outside the secondary containment. The spill did not contain any detectable levels of beryllium. However, original sampling data from the 231A tank indicated levels of 0.2 to 0.3 micrograms per liter (ug/liter) of beryllium. Therefore, containment was established and recovery actions were developed for cleanup operations.

The spill was immediately contained and a Recovery Plan was initiated for cleanup operations. The Colorado Department of Public Health and Environment (CDPHE) was notified of the spill and the Recovery Actions that were initiated.

Impact on Environment, Safety, and Health: The assessment of actual or potential hazards to human health or the environment was determined to be minimal. The RCRA Contingency Plan was implemented due to exceeding a reportable quantity for an F-listed waste (that is, 10 pounds). However, the analytical results of the sludge samples indicated the actual levels of contamination of concern were very low and posed a minimal risk to the environment. The analytical results for all of the RCRA-regulated metals were well below the regulatory limits, with cadmium results being the highest at 104 micrograms per liter (μ g/L) (parts per billion). The radiological test results indicated all isotopes tested for were in the range of picocuries per gram concentrations, with the isotope plutonium- 241 having the highest activity at 1,808 picocuries per gram (pCi/g). The highest contamination levels found during the initial release response were in the range of 100 to 600 dpm/100 cm².









































